

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
1 September 2005 (01.09.2005)

PCT

(10) International Publication Number
WO 2005/080360 A1

(51) International Patent Classification⁷: **C07D 277/46, 417/12, 413/12, 405/12, 231/40, 285/12, 213/75, 241/20, 263/48, 417/04, 403/12, A61K 31/426, A61P 3/04, 3/06**

(21) International Application Number:
PCT/GB2005/000562

(22) International Filing Date: 15 February 2005 (15.02.2005)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
0403595.2 18 February 2004 (18.02.2004) GB
0413388.0 16 June 2004 (16.06.2004) GB

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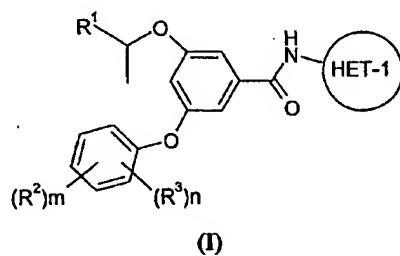
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(81) Designated States (unless otherwise indicated, for every kind of national protection available): **AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM,**

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(54) Title: COMPOUNDS



(57) Abstract: Compounds of Formula (I) wherein: R1 is methyl; R2 is selected from -C (O) NR4R5, SO2NR4R5, S (O) pR4 and HET-2; HET-1 is a 5- or 6-membered, optionally substituted C-linked heteroaryl ring; HET-2 is a 4-, 5- or 6-membered, C- or N-linked optionally substituted heterocycl ring; R3 is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano; R4 is selected from for example hydrogen, optionally substituted (1-4C) alkyl and HET-2; R5 is hydrogen or (1-4C) alkyl; or R4 and R5 together with the nitrogen atom to which they are attached may form a heterocycl ring system as defined by HET-3; HET-3 is for example an optionally substituted N-linked, 4, 5 or 6 membered, saturated or partially unsaturated heterocycl ring; p is (independently at each occurrence) 0, 1 or 2; m is 0 or 1; n is 0, 1 or 2; provided that when m is 0, then n is 1 or 2; or a salt, pro drug or solvate thereof, are described. Their use as GLK activators, pharmaceutical compositions containing them and processes for their preparation are also described.

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TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM,
ZW.

SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN,
GQ, GW, ML, MR, NE, SN, TD, TG).

(84) **Designated States (unless otherwise indicated, for every kind of regional protection available):** ARIPO (BW, GH,
GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,
ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM),
European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI,
FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO,

Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

COMPOUNDS

The present invention relates to a group of benzoyl amino heterocycll compounds which are useful in the treatment or prevention of a disease or medical condition mediated through glucokinase (GLK or GK), leading to a decreased glucose threshold for insulin secretion. In addition the compounds are predicted to lower blood glucose by increasing hepatic glucose uptake. Such compounds may have utility in the treatment of Type 2 diabetes and obesity. The invention also relates to pharmaceutical compositions comprising said compounds and to methods of treatment of diseases mediated by GLK using said compounds.

In the pancreatic β -cell and liver parenchymal cells the main plasma membrane glucose transporter is GLUT2. Under physiological glucose concentrations the rate at which GLUT2 transports glucose across the membrane is not rate limiting to the overall rate of glucose uptake in these cells. The rate of glucose uptake is limited by the rate of phosphorylation of glucose to glucose-6-phosphate (G-6-P) which is catalysed by glucokinase (GLK) [1]. GLK has a high (6-10mM) Km for glucose and is not inhibited by physiological concentrations of G-6-P [1]. GLK expression is limited to a few tissues and cell types, most notably pancreatic β -cells and liver cells (hepatocytes) [1]. In these cells GLK activity is rate limiting for glucose utilisation and therefore regulates the extent of glucose induced insulin secretion and hepatic glycogen synthesis. These processes are critical in the maintenance of whole body glucose homeostasis and both are dysfunctional in diabetes [2].

In one sub-type of diabetes, Maturity-Onset Diabetes of the Young Type 2 (MODY-2), the diabetes is caused by GLK loss of function mutations [3, 4]. Hyperglycaemia in MODY-2 patients results from defective glucose utilisation in both the pancreas and liver [5]. Defective glucose utilisation in the pancreas of MODY-2 patients results in a raised threshold for glucose stimulated insulin secretion. Conversely, rare activating mutations of GLK reduce this threshold resulting in familial hyperinsulinism [6, 6a, 7]. In addition to the reduced GLK activity observed in MODY-2 diabetics, hepatic glucokinase activity is also decreased in Type 2 diabetics [8]. Importantly, global or liver selective overexpression of GLK prevents or reverses the development of the diabetic phenotype in both dietary and genetic models of the disease [9-12]. Moreover, acute treatment of Type 2 diabetics with fructose improves glucose tolerance through stimulation of hepatic glucose utilisation [13]. This effect is believed to be

mediated through a fructose induced increase in cytosolic GLK activity in the hepatocyte by the mechanism described below [13].

Hepatic GLK activity is inhibited through association with GLK regulatory protein (GLKRP). The GLK/GLKRP complex is stabilised by fructose-6-phosphate (F6P) binding to 5 the GLKRP and destabilised by displacement of this sugar phosphate by fructose-1-phosphate (F1P). F1P is generated by fructokinase mediated phosphorylation of dietary fructose. Consequently, GLK/GLKRP complex integrity and hepatic GLK activity is regulated in a nutritionally dependent manner as F6P is dominant in the post-absorptive state whereas F1P predominates in the post-prandial state. In contrast to the hepatocyte, the pancreatic β -cell 10 expresses GLK in the absence of GLKRP. Therefore, β -cell GLK activity is regulated extensively by the availability of its substrate, glucose. Small molecules may activate GLK either directly or through destabilising the GLK/GLKRP complex. The former class of compounds are predicted to stimulate glucose utilisation in both the liver and the pancreas whereas the latter are predicted to act exclusively in the liver. However, compounds with 15 either profile are predicted to be of therapeutic benefit in treating Type 2 diabetes as this disease is characterised by defective glucose utilisation in both tissues.

GLK, GLKRP and the K_{ATP} channel are expressed in neurones of the hypothalamus, a region of the brain that is important in the regulation of energy balance and the control of food intake [14-18]. These neurones have been shown to express orectic and anorectic 20 neuropeptides [15, 19, 20] and have been assumed to be the glucose-sensing neurones within the hypothalamus that are either inhibited or excited by changes in ambient glucose concentrations [17, 19, 21, 22]. The ability of these neurones to sense changes in glucose levels is defective in a variety of genetic and experimentally induced models of obesity [23- 28]. Intracerebroventricular (icv) infusion of glucose analogues, that are competitive 25 inhibitors of glucokinase, stimulate food intake in lean rats [29, 30]. In contrast, icv infusion of glucose suppresses feeding [31]. Thus, small molecule activators of GLK may decrease food intake and weight gain through central effects on GLK. Therefore, GLK activators may be of therapeutic use in treating eating disorders, including obesity, in addition to diabetes. The hypothalamic effects will be additive or synergistic to the effects of the same compounds 30 acting in the liver and/or pancreas in normalising glucose homeostasis, for the treatment of Type 2 diabetes. Thus the GLK/GLKRP system can be described as a potential "Diabesity" target (of benefit in both Diabetes and Obesity).

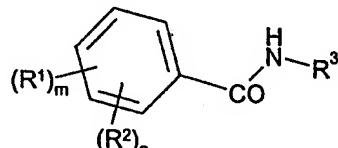
GLK is also expressed in specific entero-endocrine cells where it is believed to control the glucose sensitive secretion of the incretin peptides GIP (glucose-dependent insulinotropic polypeptide) and GLP-1 (Glucagon-Like Peptide-1) from gut K-cells and L-cells respectively (32, 33, 34). Therefore, small molecule activators of GLK may have additional beneficial effects on insulin secretion, β -cell function and survival and body weight as a consequence of stimulating GIP and GLP-1 secretion from these entero-endocrine cells.

In WO00/58293 and WO01/44216 (Roche), a series of benzylcarbamoyl compounds are described as glucokinase activators. The mechanism by which such compounds activate GLK is assessed by measuring the direct effect of such compounds in an assay in which GLK activity is linked to NADH production, which in turn is measured optically - see details of the *in vitro* assay described hereinafter. Compounds of the present invention may activate GLK directly or may activate GLK by inhibiting the interaction of GLKRP with GLK.

Further GLK activators have been described in WO03/095438 (substituted phenylacetamides, Roche), WO03/055482 (carboxamide and sulphonamide derivatives, Novo Nordisk), WO2004/002481 (arylcarbonyl derivatives, Novo Nordisk), and in WO03/080585 (amino-substituted benzoylaminoheterocycles, Banyu).

Our International application Number: WO03/000267 describes a group of benzoyl amino pyridyl carboxylic acids which are activators of the enzyme glucokinase (GLK).

Our International application Number: WO03/015774 describes compounds of the Formula (A):



(A)

wherein R³ is a substituted heterocycle other than a carboxylic acid substituted pyridyl.

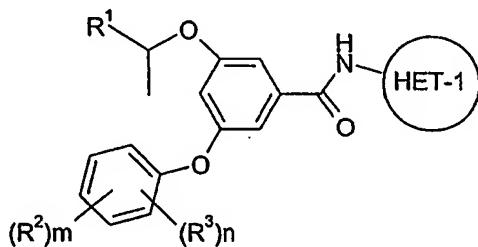
International application WO2004/076420 (Banyu) describes compounds which are generally a subset of those described in WO03/015774, wherein for example R¹ is an (substituted) alkyl ether and R² is (substituted) phenoxy.

We have surprisingly found a small group of compounds, generally a selected subgroup of those described in WO 03/015774, which have generally superior potency for the GLK enzyme, and more advantageous physical properties, including, for example, one or more of higher aqueous solubility, higher permeability, and/or lower plasma protein binding.

Consequently, such compounds having a balance of these properties would be expected to display higher plasma free drug levels and superior in vivo efficacy after oral dosing as determined, for example, by activity in Oral Glucose Tolerance Tests (OGTTs). Therefore this group of compounds would be expected to provide superior oral exposure at a lower dose and

5 thereby be particularly suitable for use in the treatment or prevention of a disease or medical condition mediated through GLK.

Thus, according to the first aspect of the invention there is provided a compound of Formula (I):



10

wherein:

R¹ is methyl;

R² is selected from -C(O)NR⁴R⁵, -SO₂NR⁴R⁵, -S(O)_pR⁴ and HET-2;

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1 or 2 further ring heteroatoms independently selected from O, N and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocyclyl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to a S(O) or S(O)₂ group, which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and

25 cyano;

R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted

with 1 group selected from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or (1-4C)alkyl;

or R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a

5 heterocycl ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl and HET-4;

R⁷ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-

10 4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-3 is an N-linked, 4, 5 or 6 membered, saturated or partially unsaturated heterocycl ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom)

independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂

15 group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an N-linked, 7 membered, saturated or partially unsaturated heterocycl ring, optionally containing 1 further heteroatom (in addition to the linking N atom) independently selected from O, S and N, wherein a -CH₂- group can optionally be replaced by a -C(O)-

20 group and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an 6-10 membered bicyclic saturated or partially unsaturated heterocycl ring, optionally containing 1 further nitrogen atom (in addition to the linking N atom) wherein a

25 -CH₂- group can optionally be replaced by a -C(O)-; which ring is optionally substituted on an available carbon or nitrogen atom by 1 substituent selected from hydroxy and R³;

R⁸ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkylamino, di(1-4C)alkylamino, HET-3 (wherein said ring is unsubstituted), (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

30 HET-4 is a 5- or 6-membered, C-or N- linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

p is (independently at each occurrence) 0, 1 or 2;

m is 0 or 1;

n is 0, 1 or 2;

provided that when m is 0, then n is 1 or 2;

or a salt, pro-drug or solvate thereof.

5 In a further aspect of the invention there is provided a compound of formula (I), or a salt, pro-drug or solvate thereof as hereinbefore defined, with the proviso that compounds exemplified in WO2004/076420, which would otherwise fall within the scope of this invention, are excluded.

In another aspect of the invention, there is provided a compound of the formula (I) as
10 hereinbefore defined, wherein

R¹ is methyl;

R² is selected from -C(O)-HET-3 and -SO₂-HET-3;

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1 or 2 further ring heteroatoms independently selected from O, N and

15 S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocyclil ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally

20 be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to a S(O) or S(O)₂ group, which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano;

25 R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or (1-4C)alkyl; or

30 R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a heterocyclil ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl and HET-4;

R⁷ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-

5 5) alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-3 is an N-linked, 4, 5 or 6 membered, saturated or partially unsaturated heterocyclyl ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom)

independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂

10 group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an N-linked, 7 membered, saturated or partially unsaturated heterocyclyl ring, optionally containing 1 further heteroatom (in addition to the linking N atom) independently selected from O, S and N, wherein a -CH₂- group can optionally be replaced by a -C(O)-

15 group and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an 6-10 membered bicyclic saturated or partially unsaturated heterocyclyl ring, optionally containing 1 further nitrogen atom (in addition to the linking N atom) wherein a

20 -CH₂- group can optionally be replaced by a -C(O)-; which ring is optionally substituted on an available carbon or nitrogen atom by 1 substituent selected from hydroxy and R³;

R⁸ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkylamino, di(1-4C)alkylamino, HET-3 (wherein said ring is unsubstituted), (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

25 HET-4 is a 5- or 6-membered, C-or N- linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

p is (independently at each occurrence) 0, 1 or 2;

m is 0 or 1;

n is 0, 1 or 2;

30 provided that when m is 0, then n is 1 or 2;

In a further aspect of the invention there is provided a compound of the formula (I) as hereinbefore defined, or a salt, pro-drug or solvate thereof, wherein:

HET-3 is an N-linked, 4 to 6 membered, saturated or partially unsaturated heterocyclyl ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom) independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸.

In another aspect of the invention, there is provided a compounds of the formula (I) as hereinbefore defined, wherein

R¹ is methyl;

10 R² is selected from -C(O)NR⁴¹R⁵¹, -SO₂NR⁴¹R⁵¹ and -S(O)_pR⁴¹;

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1 or 2 further ring heteroatoms independently selected from O, N and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from

15 R⁶;

HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocyclyl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to a S(O) or S(O)₂ group, which ring is optionally substituted on an available

20 carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano;

R⁴¹ is selected from (1-4C)alkyl [substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected

25 from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵¹ is hydrogen or (1-4C)alkyl;

R⁴ is selected from (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group

30 selected from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or (1-4C)alkyl;

or R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a heterocyclyl ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl,

5 di(1-4C)alkylamino(1-4C)alkyl and HET-4;

R⁷ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-3 is an N-linked, 4, 5 or 6 membered, saturated or partially unsaturated heterocyclyl ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom)

10 independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an N-linked, 7 membered, saturated or partially unsaturated heterocyclyl ring,

15 optionally containing 1 further heteroatom (in addition to the linking N atom) independently selected from O, S and N, wherein a -CH₂- group can optionally be replaced by a -C(O)- group and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

20 HET-3 is an 6-10 membered bicyclic saturated or partially unsaturated heterocyclyl ring, optionally containing 1 further nitrogen atom (in addition to the linking N atom) wherein a -CH₂- group can optionally be replaced by a -C(O)-; which ring is optionally substituted on an available carbon or nitrogen atom by 1 substituent selected from hydroxy and R³;

R⁸ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkylamino,

25 di(1-4C)alkylamino, HET-3 (wherein said ring is unsubstituted), (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-4 is a 5- or 6-membered, C-or N- linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

p is (independently at each occurrence) 0, 1 or 2;

30 m is 0 or 1;

n is 0, 1 or 2;

provided that when m is 0, then n is 1 or 2;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention there is provided a compound of the formula (I) as hereinbefore defined, or a salt, pro-drug or solvate thereof, wherein:

- 5 R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵], and HET-2; HET-3 as an 6-10 membered bicyclic saturated or partially unsaturated heterocycl ring, optionally containing 1 further nitrogen atom (in addition to the linking N atom) wherein a
- 10 -CH₂- group can optionally be replaced by a -C(O)-, is optionally substituted on an available carbon or nitrogen atom by 1 substituent selected from R³.

In another aspect of the invention, there is provided a compounds of the formula (I) as hereinbefore defined, wherein

R¹ is methyl;

- 15 R² is HET-2;

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1 or 2 further ring heteroatoms independently selected from O, N and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from

- 20 R⁶;

HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocycl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to a S(O) or S(O)₂ group, which ring is optionally substituted on an available

- 25 carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano;

- 30 R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or (1-4C)alkyl;

or R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a heterocyclyl ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl,

5 di(1-4C)alkylamino(1-4C)alkyl and HET-4;

R⁷ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-3 is an N-linked, 4, 5 or 6 membered, saturated or partially unsaturated heterocyclyl ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom)

10 independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an N-linked, 7 membered, saturated or partially unsaturated heterocyclyl ring,

15 optionally containing 1 further heteroatom (in addition to the linking N atom) independently selected from O, S and N, wherein a -CH₂- group can optionally be replaced by a -C(O)- group and wherein a sulphur atom in the ring may optionally be oxidised to a S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

20 HET-3 is an 6-10 membered bicyclic saturated or partially unsaturated heterocyclyl ring, optionally containing 1 further nitrogen atom (in addition to the linking N atom) wherein a -CH₂- group can optionally be replaced by a -C(O)-; which ring is optionally substituted on an available carbon or nitrogen atom by 1 substituent selected from hydroxy and R³;

R⁸ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkylamino,

25 di(1-4C)alkylamino, HET-3 (wherein said ring is unsubstituted), (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-4 is a 5- or 6-membered, C- or N-linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

p is (independently at each occurrence) 0, 1 or 2;

30 m is 0 or 1;

n is 0, 1 or 2;

provided that when m is 0, then n is 1 or 2;

or a salt, pro-drug or solvate thereof.

It will be understood that when R⁴ is -C(O)NR⁵R⁵, each R⁵ is independently selected from hydrogen and (1-4C)alkyl, and therefore this definition of R⁴ includes (but is not limited to) -CONH₂, -CONHMe, -CONMe₂ and -CONMeEt.

5 It will be understood that where a compound of the formula (I) contains more than one HET-2 ring, they may be the same or different.

It will be understood that where a compound of the formula (I) contains more than one group R⁴, they may be the same or different.

10 It will be understood that where a compound of the formula (I) contains more than one group R⁵, they may be the same or different.

It will be understood that where a compound of the formula (I) contains more than one group R⁸, they may be the same or different.

A similar convention applies for all other groups and substituents on a compound of formula (I) as hereinbefore defined.

15 Compounds of Formula (I) may form salts which are within the ambit of the invention. Pharmaceutically acceptable salts are preferred although other salts may be useful in, for example, isolating or purifying compounds.

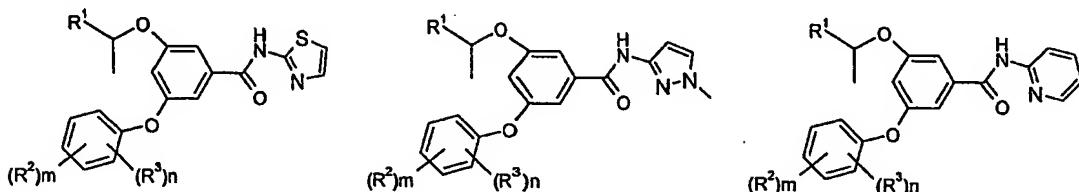
In another aspect, the invention relates to compounds of formula (I) as hereinabove defined or to a pharmaceutically acceptable salt.

20 In another aspect, the invention relates to compounds of formula (I) as hereinabove defined or to a pro-drug thereof. Suitable examples of pro-drugs of compounds of formula (I) are in-vivo hydrolysable esters of compounds of formula (I). Therefore in another aspect, the invention relates to compounds of formula (I) as hereinabove defined or to an in-vivo hydrolysable ester thereof.

25 In this specification the generic term "alkyl" includes both straight-chain and branched-chain alkyl groups. However references to individual alkyl groups such as "propyl" are specific for the straight chain version only and references to individual branched-chain alkyl groups such as *t*-butyl are specific for the branched chain version only. For example, "(1-4C)alkyl" includes methyl, ethyl, propyl, isopropyl and *t*-butyl. An analogous convention 30 applies to other generic terms.

For the avoidance of doubt, reference to the group HET-1 containing a nitrogen in the 2-position, is intended to refer to the 2-position relative to the amide nitrogen atom to which

the group is attached. For example, the following structures are encompassed (but not limited to):



Suitable examples of HET-1 as a 5- or 6-membered, C-linked heteroaryl ring as

5 hereinbefore defined, include thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, tetrazolyl and triazolyl.

It will be understood that HET-2 can be a saturated, or partially or fully unsaturated ring.

10 Suitable examples of HET-2 include azetidinyl, furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, morpholino, morpholinyl, piperidinyl, piperazinyl, morpholinyl, thiomorpholinyl, pyrrolyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oxoimidazolidinyl, 2,4-dioxoimidazolidinyl, 2-oxo-1,3,4-(4-triazolinyl), 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyrananyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 1,2,4-triazolyl, 1,2,3-triazolyl, pyranyl, and 4-pyridonyl.

15

It will be understood that HET-2 may be linked by any appropriate available C or N atom, therefore for example, for HET-2 as "imidazolyl" includes 1-, 2-, 4- and 5-imidazolyl.

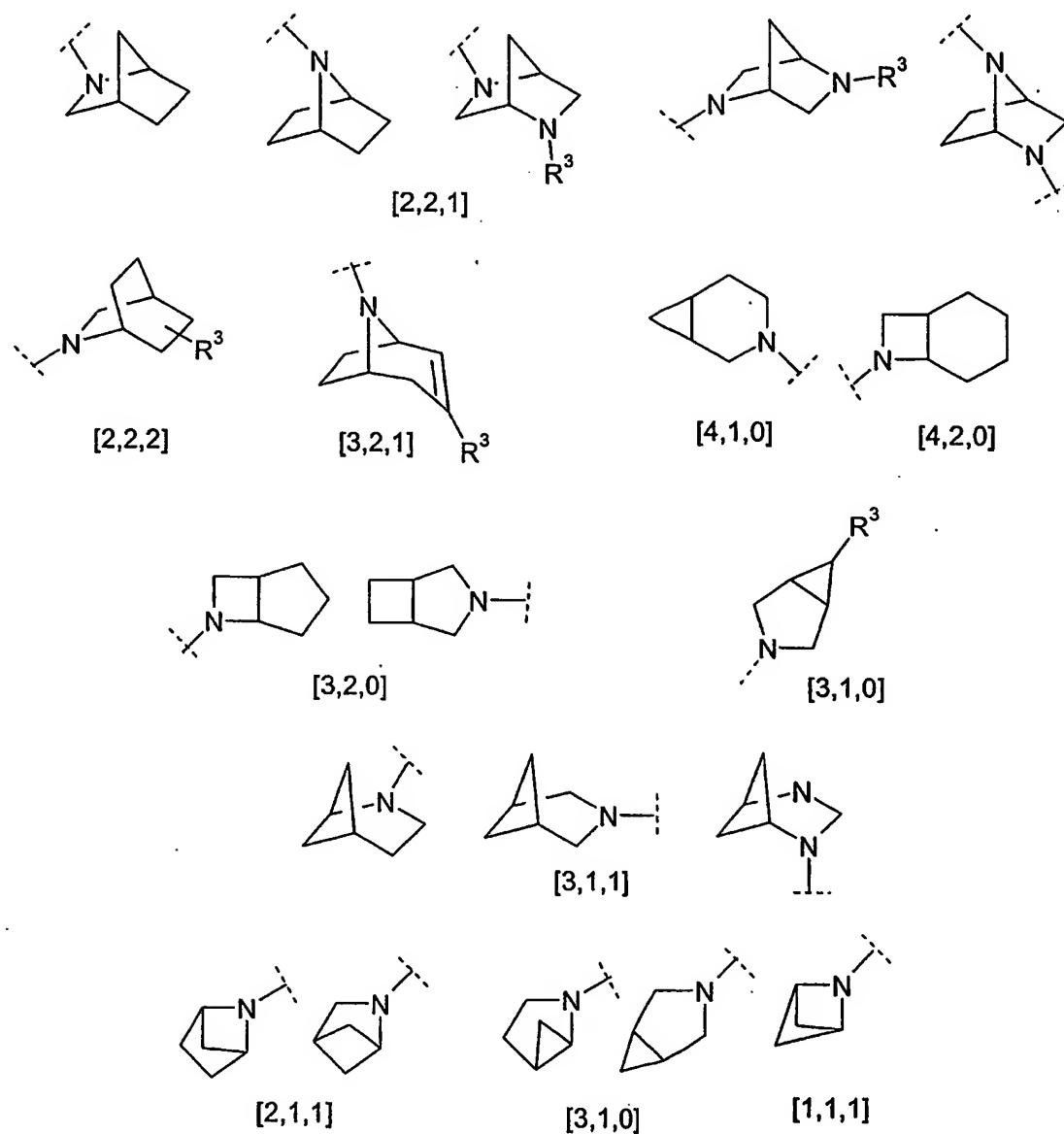
20 Suitable examples of HET-3 as a 4-6 membered saturated or partially unsaturated heterocyclic ring are morpholino, piperidinyl, piperazinyl, pyrrolidinyl and azetidinyl.

Suitable examples of HET-3 as a 7-membered saturated or partially unsaturated heterocyclic ring are homopiperazinyl, homo-morpholino, homo-thiomorpholino (and versions thereof wherein the sulfur is oxidised to an SO or S(O)₂ group) and homo-

25 piperidinyl.

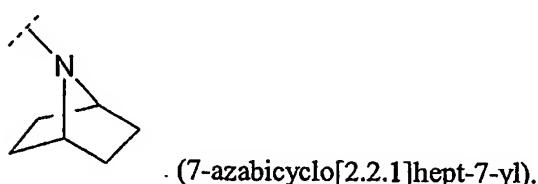
Suitable examples of HET-3 as an 6-10 membered bicyclic heterocyclic ring are bicyclic saturated or partially unsaturated heterocyclic ring such as those illustrated by the structures shown below (wherein the dotted line indicates the point of attachment to the rest of the molecule):

- 14 -



In particular HET-3 is a [2,2,1] system such as

5



Suitable examples of HET-3 are morpholino, piperidinyl, piperazinyl, pyrrolidinyl and azetidinyl.

Suitable examples of HET-4 are furyl, pyrrolyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl and triazolyl.

It will be appreciated that, where definitions of heterocyclic groups HET-1 to HET-4 encompass heteroaryl rings which may be substituted on nitrogen, such substitution may not result in charged quaternary nitrogen atoms. It will be appreciated that the definitions of HET-1 to HET-4 are not intended to include any O-O, O-S or S-S bonds. It will be appreciated that the definitions of HET-1 to HET-4 are not intended to include unstable structures.

Examples of (1-4C)alkyl include methyl, ethyl, propyl, isopropyl, butyl and tert-butyl; examples of (3-6C)cycloalkyl include cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl; examples of halo include fluoro, chloro, bromo and iodo; examples of hydroxy(1-4C)alkyl include hydroxymethyl, 1-hydroxyethyl, 2-hydroxyethyl, 2-hydroxypropyl, 3-hydroxypropyl, 1-hydroxyisopropyl and 4-hydroxybutyl; examples of (1-4C)alkoxy(1-4C)alkyl include methoxymethyl, ethoxymethyl, tert-butoxymethyl, 2-methoxyethyl, 2-ethoxyethyl, 15 methoxypropyl, 2-methoxypropyl and methoxybutyl; examples of (1-4C)alkylS(O)p(1-4C)alkyl include methylsulfinylmethyl, ethylsulfinylmethyl, ethylsulfinylethyl, methylsulfinylpropyl, methylsulfinylbutyl, methylsulfonylmethyl, ethylsulfonylmethyl, ethylsulfonylethyl, methylsulfonylpropyl, methylsulfonylbutyl, methylthiomethyl, ethylthiomethyl, ethylthioethyl, methylthiopropyl, and methylthiobutyl; examples of amino(1-20 4C)alkyl include aminomethyl, aminoethyl, 2-aminopropyl, 3-aminopropyl, 1-aminoisopropyl and 4-aminobutyl; examples of (1-4C)alkylamino(1-4C)alkyl include (N-methyl)aminomethyl, (N-ethyl)aminomethyl, 1-((N-methyl)amino)ethyl, 2-((N-methyl)amino)ethyl, (N-ethyl)aminoethyl, (N-methyl)aminopropyl, and 4-((N-methyl)amino)butyl; examples of di(1-4C)alkylamino(1-4C)alkyl include 25 dimethylaminomethyl, methyl(ethyl)aminomethyl, methyl(ethyl)aminoethyl, (N,N-diethyl)aminoethyl, (N,N-dimethyl)aminopropyl and (N,N-dimethyl)aminobutyl; examples of (1-4C)alkylamino include methylamino, ethylamino, propylamino, isopropylamino, butylamino and tert-butylamino; examples of di(1-4C)alkylamino include dimethylamino, methyl(ethyl)amino, diethylamino, dipropylamino, di-isopropylamino and dibutylamino; 30 examples of -C(O)(1-4C)alkyl include methylcarbonyl, ethylcarbonyl, propylcarbonyl and tert-butyl carbonyl.

It is to be understood that, insofar as certain of the compounds of Formula (I) defined above may exist in optically active or racemic forms by virtue of one or more asymmetric carbon atoms, the invention includes in its definition any such optically active or racemic form which possesses the property of stimulating GLK directly or inhibiting the GLK/GLKRP interaction. The synthesis of optically active forms may be carried out by standard techniques of organic chemistry well known in the art, for example by synthesis from optically active starting materials or by resolution of a racemic form. It is also to be understood that certain compounds may exist in tautomeric forms and that the invention also relates to any and all tautomeric forms of the compounds of the invention which activate GLK.

10 In one embodiment of the invention are provided compounds of formula (I), in an alternative embodiment are provided pharmaceutically-acceptable salts of compounds of formula (I), in a further alternative embodiment are provided in-vivo hydrolysable esters of compounds of formula (I), and in a further alternative embodiment are provided pharmaceutically-acceptable salts of in-vivo hydrolysable esters of compounds of formula (I).

15 Particular values of variable groups are as follows. Such values may be used where appropriate with any of the values, definitions, claims, aspects or embodiments defined hereinbefore or hereinafter. In particular, each may be used as an individual limitation on the broadest definition of formula (I). Further, each of the following values may be used in combination with one or more of the other following values to limit the broadest definition of formula (I).

(1) R² is -C(O)NR⁴R⁵

(2) R² is -SO₂NR⁴R⁵

(3) R² is -S(O)_pR⁴

20 (4) R² is HET-2

(5) m is 1 and R² is in the para position relative to the ether linkage

(6) m is 1 and n is 0 or 1

(7) m is 1 and n is 0

(8) m is 1 and n is 1

25 (9) m is 1, n is 0 and R² is in the para position relative to the ether linkage

(10) m is 1, n is 1, R² is in the para position relative to the ether linkage, R³ is in the ortho position relative to the ether linkage

(11) m is 1, n is 1, R² is in the para position relative to the ether linkage, R³ is in the meta position relative to the ether linkage

(12) n is 0

(13) n is 1

5 (14) n is 2

(15) n is 2 and both R³ are halo

(16) n is 2 and each R³ is independently halo or methoxy

(17) m is 1, n is 2 and R² is in the para position relative to the ether linkage

(18) m is 1, n is 2, R² is in the para position relative to the ether linkage and each R³ is in an ortho position relative to the ether linkage

10 (19) m is 1, n is 2, both R³ are halo, R² is in the para position relative to the ether linkage and each R³ is in an ortho position relative to the ether linkage

(20) R³ is fluoromethyl or difluoromethyl

(21) R³ is halo or trifluoromethyl

15 (22) R³ is halo

(23) R³ is chloro or fluoro

(24) R³ is fluoro

(25) R³ is methoxy

(26) n is 2 and both R³ are fluoro,

20 (27) n is 2, both R³ are fluoro and are in the 3- and 5-positions (meta-positions) relative to the ether linkage

(28) m is 1, n is 2, R² is in the para position relative to the ether linkage, both R³ are fluoro and are in the 3- and 5-positions relative to the ether linkage

(29) p is 0

25 (30) p is 1

(31) p is 2

(32) HET-1 is a 5-membered heteroaryl ring

(33) HET-1 is a 6-membered heteroaryl ring

(34) HET-1 is substituted with 1 or 2 substituents independently selected from R⁶

30 (35) HET-1 is substituted with 1 substituent selected from R⁶

(36) HET-1 is unsubstituted

(37) HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, and triazolyl

(38) HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, 5 oxazolyl, isoxazolyl and oxadiazolyl

(39) HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl

(40) HET-1 is selected from thiazolyl, pyrazolyl and oxazolyl

(41) HET-1 is selected from thiadiazolyl and oxadiazolyl

(42) HET-1 is selected from 1,3,4-thiadiazolyl and 1,3,4-oxadiazolyl

10 (43) HET-1 is selected from 1,2,4-oxadiazolyl and 1,2,4-oxadiazolyl

(44) HET-1 is pyrazolyl

(45) HET-1 is pyridyl or pyrazinyl

(46) HET-1 is selected from thiazolyl, pyrazolyl, thiadiazolyl and pyridyl;

(47) R⁶ is selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, di(1-4C)alkylamino(1-15 4C)alkyl and HET-4

(48) R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, hydroxymethyl, methoxymethyl, aminomethyl, N-methylaminomethyl, dimethylaminomethyl

(49) R⁶ is selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, and di(1-20 4C)alkylamino(1-4C)alkyl

(50) R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl

(51) R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, hydroxymethyl and methoxymethyl

25 (52) R⁶ is selected from methyl, ethyl, bromo, chloro and fluoro

(53) R⁶ is methyl

(54) R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, dimethylaminomethyl, hydroxymethyl and methoxymethyl

(55) R⁶ is selected from methyl, ethyl, aminomethyl, N-methylaminomethyl,

30 dimethylaminomethyl, hydroxymethyl and methoxymethyl

(56) R⁶ is selected from methyl, ethyl, isopropyl and methoxymethyl

(57) when 2 substituents R⁶ are present, both are selected from methyl, ethyl, bromo, chloro and fluoro; preferably both are methyl

(58) R⁶ is selected from (1-4C)alkylS(O)p(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl and HET-4

(59) R⁶ is HET-4

(60) HET-4 is selected from furyl, pyrrolyl and thieryl

5 (61) HET-4 is furyl

(62) R⁴ is hydrogen

(63) R⁴ is (1-4C)alkyl [substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵]

10 (64) R⁴ is (1-4C)alkyl [substituted by 1 substituent selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl and -C(O)NR⁵R⁵]

(65) R⁴ is (1-4C)alkyl

(66) R⁴ is (1-4C)alkyl substituted by -OR⁵

(67) R⁴ is (1-4C)alkyl substituted by HET-2

15 (68) R⁴ is (3-6C)cycloalkyl, particularly cyclopropyl

(69) R⁴ is (3-6C)cycloalkyl substituted by a group selected from R⁷

(70) R⁴ is (3-6C)cycloalkyl substituted by a group selected from -OR⁵ and (1-4C)alkyl

(71) R⁴ is HET-2

(72) R⁴ is selected from hydrogen, (1-4C)alkyl, and (1-4C)alkyl substituted with -OR⁵

20 (73) HET-2 is unsubstituted

(74) HET-2 is substituted with 1 or 2 substituents independently selected from (1-4C)alkyl, hydroxy and (1-4C)alkoxy

(75) HET-2 is a fully saturated ring system

(76) HET-2 is a fully unsaturated ring system

25 (77) HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-dioxoimidazolidinyl, pyranyl and 4-pyridonyl

30 (78) HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, pyrrolidinyl, thiomorpholinyl, tetrahydrofuranyl, and tetrahydropyranyl

(79) HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl

(80) HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, 5 imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, pyrrolidonyl, 2-oxazolidinonyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl

(81) HET-2 is selected from morpholino, furyl, imidazolyl, oxazolyl, isoxazolyl, oxadiazolyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, 2-pyrrolidonyl, 2-oxazolidinonyl, 10 tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl

(82) HET-2 is selected from morpholino, furyl, imidazolyl, isoxazolyl, oxadiazolyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, 2-pyrrolidonyl, tetrahydropyranyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl

(83) R⁵ is hydrogen

15 (84) R⁵ is (1-4)alkyl, preferably methyl

(85) R⁵ is hydrogen or methyl

(86) R⁷ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, and hydroxy(1-4C)alkyl

(87) R⁷ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, and hydroxy(1-20 4C)alkyl

(88) R⁷ is selected from hydroxy, methoxy, -COMe, -CONH₂, -CONHMe, -CONMe₂, and hydroxymethyl

(89) R⁷ is selected from (1-4C)alkyl, hydroxy and (1-4C)alkoxy

(90) R⁷ is selected from methyl, ethyl, methoxy and hydroxy

25 (91) R⁷ is methyl

(92) R⁸ is selected from methyl, hydroxy, methoxy, -COMe, -CONH₂, -CONHMe, -CONMe₂, hydroxymethyl, hydroxyethyl, -NHMe and -NMe₂

(93) R⁸ is selected from morpholino, piperidinyl, piperazinyl, pyrrolidinyl and azetidinyl

(94) R⁸ is selected from methyl, -COMe, -CONH₂, hydroxyethyl and hydroxy

30 (95) R⁸ is methyl

(96) HET-3 is a fully saturated ring

(97) HET-3 is selected from morpholino, piperidinyl, piperazinyl, pyrrolidinyl and azetidinyl

(98) R⁴ and R⁵ together with the nitrogen to which they are attached form a ring as defined by HET-3

(99) HET-3 is selected from pyrrolidinyl and azetidinyl

(100) HET-3 is azetidinyl

5 (101) HET-3 is a 4, 5 or 6-membered saturated or partially unsaturated heterocyclic ring as hereinbefore defined

(102) HET-3 is a 7-membered saturated or partially unsaturated heterocyclic ring as hereinbefore defined

(103) HET-3 is an 6 to 10-membered bicyclic saturated or partially unsaturated heterocyclic

10 ring as hereinbefore defined

(104) HET-3 is 7-azabicyclo[2.2.1]hept-7-yl

(105) HET-3 is selected from morpholino, piperidinyl, piperazinyl, pyrrolidinyl, azetidinyl and 7-azabicyclo[2.2.1]hept-7-yl

(106) HET-3 is selected from piperidinyl, pyrrolidinyl, azetidinyl and 7-azabicyclo[2.2.1]hept-

15 7-yl

According to a further feature of the invention there is provided the following preferred groups of compounds of the invention:

In a futher aspect of the invention there is provided a compound of Formula (I)

20 wherein:

R¹ is methyl;

R² is selected from -C(O)NR⁴R⁵, -SO₂NR⁴R⁵, -S(O)_pR⁴ and HET-2;

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1, 2 or 3 further ring heteroatoms independently selected from O, N

25 and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

HET-2 is a 5- or 6-membered, C- or N-linked heterocyclyl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally

30 be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to an S(O) or S(O)₂ group, which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano;

R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by -OR⁵] and HET-2;

R⁵ is hydrogen or (1-4C)alkyl;

5 or R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a 4-6 membered heterocyclyl ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl and HET-4;

10 R⁷ is selected from -OR⁵ and (1-4C)alkyl;

HET-3 is an N-linked, 4 to 6 membered, saturated or partially unsaturated heterocyclyl ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom) independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein sulphur atoms in the ring may optionally be oxidised to S(O) or S(O)₂

15 groups; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸;

R⁸ is selected from -OR⁵ and (1-4C)alkyl;

HET-4 is a 5- or 6-membered, C-or N- linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

20 p is (independently at each occurrence) 0, 1 or 2;

m is 0 or 1;

n is 0, 1 or 2;

provided that when m is 0, then n is 1 or 2;

or a salt, pro-drug or solvate thereof.

25

In a further aspect of the invention there is provided a compound of Formula (I)

wherein:

R¹ is methyl;

R² is selected from -C(O)NR⁴R⁵, -SO₂NR⁴R⁵, -S(O)_pR⁴ and HET-2;

30 HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1, 2 or 3 further ring heteroatoms independently selected from O, N and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen

atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

HET-2 is a 5- or 6-membered, C- or N-linked heterocycl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally

5 be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to an S(O) or S(O)₂ group, which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano;

10 R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by -OR⁵] and HET-2;

R⁵ is hydrogen or (1-4C)alkyl;

or R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a 4-6 membered heterocycl ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-

15 4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl and HET-4;

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

HET-3 is an N-linked, 4 to 6 membered, saturated or partially unsaturated heterocycl ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom)

20 independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to an S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an N-linked, 7 membered, saturated or partially unsaturated heterocycl ring,

25 optionally containing 1 further heteroatom (in addition to the linking N atom) independently selected from O, S and N, wherein a -CH₂- group can optionally be replaced by a -C(O)- group and wherein a sulphur atom in the ring may optionally be oxidised to an S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

30 HET-3 is an 6-10 membered bicyclic saturated or partially unsaturated heterocycl ring, optionally containing 1 further nitrogen atom (in addition to the linking N atom), wherein a

-CH₂- group can optionally be replaced by a -C(O)-; which ring is optionally substituted on an available carbon or nitrogen atom by 1 substituent selected from R³;

R⁸ is selected from -OR⁵ and (1-4C)alkyl;

R⁸ is selected from -OR⁵ and (1-4C)alkyl;

5 HET-4 is a 5- or 6-membered, C-or N-linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

p is (independently at each occurrence) 0, 1 or 2;

m is 0 or 1;

n is 0, 1 or 2;

10 provided that when m is 0, then n is 1 or 2;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein:

15 R¹ is methyl;

R² is selected from -C(O)NR⁴R⁵, -SO₂NR⁴R⁵, -S(O)_pR⁴ and HET-2;

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1 or 2 further ring heteroatoms independently selected from O, N and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom

20 provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocyclyl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally

25 be oxidised to an S(O) or S(O)₂ group, which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano;

R⁴ is selected from (1-4C)alkyl [substituted by 1 or 2 substituents independently selected from

30 HET-2, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵];

R⁵ is hydrogen or (1-4C)alkyl;

or R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a 4-6 membered heterocyclyl ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl and HET-4;

R⁷ is selected from -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-3 is an N-linked, 4 to 6 membered, saturated or partially unsaturated heterocyclyl ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom)

10 independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to an S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸;

R⁸ is selected from -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkylamino, di(1-4C)alkylamino,

15 HET-3 (wherein said ring is unsubstituted), (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-4 is a 5- or 6-membered, C-or N-linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

p is (independently at each occurrence) 0, 1 or 2;

20 m is 0 or 1;

n is 0, 1 or 2;

provided that when m is 0, then n is 1 or 2;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as

25 hereinbefore defined wherein:

R¹ is methyl;

R² is selected from -C(O)NR⁴R⁵, -SO₂NR⁴R⁵, -S(O)_pR⁴ and HET-2;

HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1 or 2 further ring heteroatoms independently selected from O, N and

30 S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocyclyl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to an S(O) or S(O)₂ group, which ring is optionally substituted on an available

5 carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano;

R⁴ is selected from (1-4C)alkyl [substituted by 1 or 2 substituents independently selected from HET-2, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and

10 -C(O)NR⁵R⁵];

R⁵ is hydrogen or (1-4C)alkyl;

or R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a 4-6 membered heterocyclyl ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-

15 4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl and HET-4;

R⁷ is selected from -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-3 is an N-linked, 4 to 6 membered, saturated or partially unsaturated heterocyclyl ring,

20 optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom)

independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to an S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

25 HET-3 is an N-linked, 7 membered, saturated or partially unsaturated heterocyclyl ring,

optionally containing 1 further heteroatom (in addition to the linking N atom) independently selected from O, S and N, wherein a -CH₂- group can optionally be replaced by a -C(O)- group and wherein a sulphur atom in the ring may optionally be oxidised to an S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2

30 substituents independently selected from R⁸; or

HET-3 is an 6-10 membered bicyclic saturated or partially unsaturated heterocyclyl ring, optionally containing 1 further nitrogen atom (in addition to the linking N atom), wherein a

-CH₂- group can optionally be replaced by a -C(O)-; which ring is optionally substituted on an available carbon or nitrogen atom by 1 substituent selected from R³;

R⁸ is selected from -OR⁵ and (1-4C)alkyl;

R⁸ is selected from -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkylamino, di(1-4C)alkylamino,

5 HET-3 (wherein said ring is unsubstituted), (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-4 is a 5- or 6-membered, C-or N-linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

p is (independently at each occurrence) 0, 1 or 2;

10 m is 0 or 1;

n is 0, 1 or 2;

provided that when m is 0, then n is 1 or 2;

or a salt, pro-drug or solvate thereof.

15 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is a 5- or 6-membered heteroaryl ring;

20 R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl; R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵];

R⁵ is hydrogen or methyl;

25 HET-2 is a 5- or 6-membered heterocyclic ring as hereinbefore defined, containing 1 or 2 heteroatoms independently selected from O, N and S; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

30 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is a 5- or 6-membered heteroaryl ring;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from

5 HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵];

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

10 HET-2 is a 5- or 6- membered heterocycl ring as hereinbefore defined, containing 1 or 2 heteroatoms independently selected from O, N and S; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

15 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl,

20 isoxazolyl and oxadiazolyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl and -C(O)NR⁵R⁵];

25 R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-

30 dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-dioxoimidazolidinyl, pyranyl and 4-pyridonyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

- 5 R¹ is methyl;
m is 1 and n is 0 or 1;
HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;
R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;
R³ is halo or trifluoromethyl;
- 10 R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl and -C(O)NR⁵R⁵];
R⁵ is hydrogen or methyl;
R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;
- 15 HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-dioxoimidazolidinyl, pyranyl and 4-pyridonyl; and
- 20 R⁷ is selected from -OR⁵ and (1-4C)alkyl;
or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

- 25 R¹ is methyl;
m is 1 and n is 0 or 1;
HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl and oxadiazolyl;
R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;
- 30 R³ is halo or trifluoromethyl;
R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl and -C(O)NR⁵R⁵];
R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl,

5 1,2,4-triazolyl and 1,2,3-triazolyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as

10 hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

15 R³ is halo or trifluoromethyl;

R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl and -C(O)NR⁵R⁵];

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl,

20 and dimethylaminomethyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

25 or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as

hereinbefore defined wherein

R¹ is methyl;

30 m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

R⁴ is selected from hydrogen, (1-4C)alkyl, [optionally substituted by -OR⁵] and HET-2;

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl,

5 and dimethylaminomethyl;

HET-2 is selected from morpholino, furyl, imidazolyl, isoxazolyl, oxadiazolyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, 2-pyrrolidonyl, tetrahydropyranlyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

10 or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

15 m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

20 R⁴ is selected from hydrogen, (1-4C)alkyl, [optionally substituted by -OR⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

25 HET-2 is selected from morpholino, furyl, imidazolyl, isoxazolyl, oxadiazolyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, 2-pyrrolidonyl, tetrahydropyranlyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

30

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl and pyridazinyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

5 R⁴ is selected from hydrogen, (1-4C)alkyl, [optionally substituted by -OR⁵] and HET-2;

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from morpholino, furyl, imidazolyl, isoxazolyl, oxadiazolyl, piperidinyl,

10 piperazinyl, 3-oxopiperazinyl, pyrrolidinyl, 2-pyrrolidonyl, tetrahydropyranlyl, 1,1-dioxotetrahydrothienyl, and 2-oxoimidazolidinyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

15 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl

20 and oxadiazolyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

R⁴ is selected from (1-4C)alkyl, [optionally substituted by -OR⁵] and HET-2;

R⁵ is hydrogen or methyl;

25 R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from piperidinyl, piperazinyl, 3-oxopiperazinyl, 2-pyrrolidonyl,

2,5-dioxopyrrolidinyl, 2-oxotetrahydrofuranlyl, tetrahydrofuranlyl, tetrahydropyranlyl, 2-

oxoimidazolidinyl, and 2,4-dioxoimidazolidinyl; and

30 R⁷ is (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

5 HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

R⁴ is selected from (1-4C)alkyl, [optionally substituted by -OR⁵], (3-6C)cycloalkyl (optionally

10 substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from piperidinyl, piperazinyl, 3-oxopiperazinyl, 2-pyrrolidonyl,

15 2,5-dioxopyrrolidinyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranly, 2-oxoimidazolidinyl, and 2,4-dioxoimidazolidinyl; and

R⁷ is (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

20 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl

25 and oxadiazolyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

R⁴ is selected from (1-4C)alkyl, [optionally substituted by -OR⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

30 R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is piperidinyl or piperazinyl; and

R⁷ is (1-4C)alkyl;
or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as
5 hereinbefore defined wherein

R¹ is methyl;
m is 1 and n is 0;
HET-1 is selected from thiazolyl, thiadiazolyl and pyrazolyl;
R² is -CONR⁴R⁵;
10 R⁴ is piperidinyl optionally substituted with methyl;
R⁵ is hydrogen or methyl;
R⁶ is methyl;
or a salt, pro-drug or solvate thereof.

15 In a further aspect of the invention is provided a compound of the formula (I) as
hereinbefore defined wherein
R¹ is methyl;
m is 1 and n is 0 or 1;
HET-1 is selected from pyridyl and pyridazinyl;
20 R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;
R³ is halo or trifluoromethyl;
R⁴ is selected from (1-4C)alkyl, [optionally substituted by -OR⁵], (3-6C)cycloalkyl (optionally
substituted with 1 group selected from R⁷) and HET-2;
R⁵ is hydrogen or methyl;
25 R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl,
and dimethylaminomethyl;
HET-2 is selected from piperidinyl, piperazinyl, 3-oxopiperazinyl, 2-pyrrolidonyl,
2,5-dioxopyrrolidinyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl,
tetrahydropyrananyl, 2-oxoimidazolidinyl, and 2,4-dioxoimidazolidinyl; and
30 R⁷ is (1-4C)alkyl;
or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

5 HET-1 is selected from pyridyl and pyridazinyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

R⁴ is selected from (1-4C)alkyl, [optionally substituted by -OR⁵] and HET-2;

R⁵ is hydrogen or methyl;

10 R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is piperidinyl or piperazinyl; and

R⁷ is (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

15

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

20 HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is -CONR⁴R⁵ or -SO₂NR⁴R⁵;

R³ is halo or trifluoromethyl;

R⁴ and R⁵ together with the nitrogen to which they are attached form a morpholino,

25 piperidinyl, piperazinyl, pyrrolidinyl or azetidinyl ring, which ring is optionally substituted on a carbon or nitrogen atom by (1-4C)alkyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

or a salt, pro-drug or solvate thereof.

30

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl and pyridazinyl;

R² is $-\text{CONR}^4\text{R}^5$ or $-\text{SO}_2\text{NR}^4\text{R}^5$;

R³ is halo or trifluoromethyl;

5 R⁴ and R⁵ together with the nitrogen to which they are attached form a morpholino, piperidinyl, piperazinyl, pyrrolidinyl or azetidinyl ring, which ring is optionally substituted on a carbon or nitrogen atom by (1-4C)alkyl;
R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;
10 or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

15 m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is $-\text{CONR}^4\text{R}^5$ or $-\text{SO}_2\text{NR}^4\text{R}^5$;

R³ is halo or trifluoromethyl;

20 R⁴ and R⁵ together with the nitrogen to which they are attached form a morpholino, piperidinyl, piperazinyl, pyrrolidinyl or azetidinyl ring, which ring is optionally substituted on a carbon or nitrogen atom by R⁸;
R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl; and
25 R⁸ is selected from hydroxy, (1-4C)alkoxy and (1-4C)alkyl;
or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

30 m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is $-\text{CONR}^4\text{R}^5$ or $-\text{SO}_2\text{NR}^4\text{R}^5$;

R³ is halo or trifluoromethyl;

R⁴ and R⁵ together with the nitrogen to which they are attached form a morpholino, piperidinyl, piperazinyl, pyrrolidinyl or azetidinyl ring, which ring is optionally substituted on a carbon or nitrogen atom by R⁸;

5 R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl; and

R⁸ is pyrrolidine or piperidine;

or a salt, pro-drug or solvate thereof.

10 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0;

HET-1 is selected from thiazolyl, thiadiazolyl and pyrazolyl;

15 R² is -CONR⁴R⁵;

R⁴ and R⁵ together with the nitrogen to which they are attached form a piperidinyl, or piperazinyl ring, which ring is optionally substituted on a carbon or nitrogen atom by (1-4C)alkyl or by a pyrrolidinyl ring;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl,

20 and dimethylaminomethyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

25 R¹ is methyl;

m is 1 and n is 0;

HET-1 is selected from thiazolyl, thiadiazolyl and pyrazolyl;

R² is -CONR⁴R⁵;

R⁴ and R⁵ together with the nitrogen to which they are attached form an azetidinyl ring which

30 ring is optionally substituted on a carbon atom by hydroxy;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, hydroxymethyl, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0;

5 HET-1 is selected from thiazolyl, thiadiazolyl and pyrazolyl;

R² is -CONR⁴R⁵;

R⁴ and R⁵ together with the nitrogen to which they are attached form a 7-membered ring HET-3 which ring is optionally substituted on a carbon or nitrogen atom by methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, hydroxymethyl, aminomethyl, N-

10 methylaminomethyl, and dimethylaminomethyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

15 R¹ is methyl;

m is 1 and n is 0;

HET-1 is selected from thiazolyl, thiadiazolyl and pyrazolyl;

R² is -CONR⁴R⁵;

R⁴ and R⁵ together with the nitrogen to which they are attached form a 6-10 membered

20 bicyclic heterocyclic ring HET-3;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, hydroxymethyl, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

or a salt, pro-drug or solvate thereof.

25 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is a 5- or 6-membered heteroaryl ring;

30 R² is -S(O)pR⁴;

p is 1 or 2;

R³ is halo or trifluoromethyl;

R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵];

R⁵ is hydrogen or methyl;

5 HET-2 is a 5- or 6-membered heterocycl ring as hereinbefore defined, containing 1 or 2 heteroatoms independently selected from O, N and S; and
R⁷ is selected from -OR⁵ and (1-4C)alkyl;
or a salt, pro-drug or solvate thereof.

10 In a further aspect of the invention is provided a compound of the formula (I) as
hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is a 5- or 6-membered heteroaryl ring;

15 R² is -S(O)pR⁴;

p is 1 or 2;

R³ is halo or trifluoromethyl;

R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from

20 R⁷) and -C(O)NR⁵R⁵];

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl];

HET-2 is a 5- or 6-membered heterocycl ring as hereinbefore defined, containing 1 or 2

25 heteroatoms independently selected from O, N and S; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as

30 hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is -S(O)pR⁴;

p is 1 or 2;

5 R³ is halo or trifluoromethyl;

R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl and -C(O)NR⁵R⁵];

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl,

10 and dimethylaminomethyl;

HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-

15 dioxoimidazolidinyl, pyranyl and 4-pyridonyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as

20 hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

25 R² is -S(O)pR⁴;

p is 1 or 2;

R³ is halo or trifluoromethyl;

R⁴ is selected from hydrogen, (1-4C)alkyl, [optionally substituted by -OR⁵] and HET-2;

R⁵ is hydrogen or methyl;

30 R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

5 or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

10 m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is -S(O)pR⁴;

p is 1 or 2;

15 R³ is halo or trifluoromethyl;

R⁴ is selected from hydrogen, (1-4C)alkyl, [optionally substituted by -OR⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl,

20 and dimethylaminomethyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

25 or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

30 m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

R² is -S(O)pR⁴;

p is 1 or 2;

R³ is halo or trifluoromethyl;

R⁴ is (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl and -C(O)NR⁵R⁵];

R⁵ is hydrogen or methyl;

5 R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl,

10 tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-dioxoimidazolidinyl, pyranyl and 4-pyridonyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;
or a salt, pro-drug or solvate thereof.

15 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

20 R² is -S(O)pR⁴;

p is 1 or 2;

R³ is halo or trifluoromethyl;

R⁴ is selected from hydrogen, (1-4C)alkyl, [optionally substituted by -OR⁵] and HET-2;

R⁵ is hydrogen or methyl;

25 R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl; and

30 R⁷ is selected from -OR⁵ and (1-4C)alkyl;
or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

5 HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

R² is -S(O)pR⁴;

p is 1 or 2;

R³ is halo or trifluoromethyl;

R⁴ is selected from hydrogen, (1-4C)alkyl, [optionally substituted by -OR⁵], (3-6C)cycloalkyl

10 (optionally substituted with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or methyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl,

15 pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl,

1,2,4-triazolyl and 1,2,3-triazolyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

20 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl,

25 isoxazolyl and oxadiazolyl;

R² is -S(O)pR⁴;

p is 1 or 2;

R³ is halo or trifluoromethyl;

R⁴ is (1-4C)alkyl;

30 R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0;

5 HET-1 is selected from thiazolyl, thiadiazolyl and pyrazolyl;

R² is -S(O)pR⁴;

p is 1 or 2;

R⁴ is (1-4C)alkyl;

R⁶ is methyl;

10 or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

m is 1 and n is 0;

15 HET-1 is selected from thiazolyl, thiadiazolyl and pyrazolyl;

R² is -S(O)pR⁴;

p is 1 or 2;

R⁴ is (3-6C)cycloalkyl;

R⁶ is methyl;

20 or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

25 m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

R² is -S(O)pR⁴;

p is 1 or 2;

R³ is halo or trifluoromethyl;

30 R⁴ is (1-4C)alkyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

5 HET-1 is a 5- or 6-membered heteroaryl ring;

R² is HET-2;

R³ is halo or trifluoromethyl;

R⁵ is hydrogen or (1-4C)alkyl;

HET-2 is a 5- or 6- membered heterocycl ring as hereinbefore defined, containing 1 or 2

10 heteroatoms independently selected from O, N and S; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;.

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as

15 hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

20 R² is HET-2;

R³ is halo or trifluoromethyl;

R⁵ is hydrogen or methyl;

HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-

25 dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-dioxoimidazolidinyl, pyranyl and 4-pyridonyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

30

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl and oxadiazolyl;

R² is HET-2;

5 R³ is halo or trifluoromethyl;

R⁵ is hydrogen or methyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl; and

10 R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

15 R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

R² is HET-2;

R³ is halo or trifluoromethyl;

20 R⁵ is hydrogen or methyl;

HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-

25 dioxoimidazolidinyl, pyranyl and 4-pyridonyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as

30 hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

R² is HET-2;

R³ is halo or trifluoromethyl;

R⁵ is hydrogen or methyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl,

5 pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl,
1,2,4-triazolyl and 1,2,3-triazolyl; and

R⁷ is selected from -OR⁵ and (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

10 In a further aspect of the invention is provided a compound of the formula (I) as
hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl,
15 isoxazolyl and oxadiazolyl;

R² is HET-2;

R³ is halo or trifluoromethyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl,
and dimethylaminomethyl;

20 HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-
oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-
dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl,
tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-
dioxoimidazolidinyl, pyranyl and 4-pyridonyl; and

25 R⁷ is (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as
hereinbefore defined wherein

30 R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, isothiazolyl, thiadiazolyl, pyrazolyl, imidazolyl, oxazolyl,
isoxazolyl and oxadiazolyl;

R² is HET-2;

R³ is halo or trifluoromethyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

5 HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl, pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl; and

R⁷ is (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

10

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

15 HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

R² is HET-2;

R³ is halo or trifluoromethyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

20 HET-2 is selected from azetidinyl, morpholino, morpholinyl, piperidinyl, piperazinyl, 3-oxopiperazinyl, thiomorpholinyl, pyrrolidinyl, pyrrolidonyl, 2,5-dioxopyrrolidinyl, 1,1-dioxotetrahydrothienyl, 2-oxazolidinonyl, 2-oxotetrahydrofuranyl, tetrahydrofuranyl, tetrahydropyranyl, 1,1-dioxothiomorpholino, 1,3-dioxolanyl, 2-oxoimidazolidinyl, 2,4-dioxoimidazolidinyl, pyranyl and 4-pyridonyl; and

25 R⁷ is (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

30 R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from pyridyl, pyrazinyl, pyridazinyl and pyrimidinyl;

R² is HET-2;

R³ is halo or trifluoromethyl;

R⁶ is selected from methyl, ethyl, bromo, chloro, fluoro, aminomethyl, N-methylaminomethyl, and dimethylaminomethyl;

HET-2 is selected from furyl, thienyl, thiazolyl, isothiazolyl, thiadiazolyl, pyridyl, pyrazinyl,

5 pyridazinyl, pyrazolyl, imidazolyl, pyrimidinyl, oxazolyl, isoxazolyl, oxadiazolyl, pyrrolyl, 1,2,4-triazolyl and 1,2,3-triazolyl; and

R⁷ is (1-4C)alkyl;

or a salt, pro-drug or solvate thereof.

10 In a further aspect of the invention is provided a compound of the formula (I) as hereinbefore defined wherein

R¹ is methyl;

m is 1 and n is 0 or 1;

HET-1 is selected from thiazolyl, pyrazolyl, N-methylpyrazolyl, thiadiazolyl, pyridyl,

15 pyrazinyl, isoxazolyl; 5-methylisoxazolyl, furyl, dimethylaminomethylthiazolyl, and methylthiadiazolyl;

R² is selected from N-methylpiperazin-4-ylcarbonyl, 2-(aminocarbonyl)-pyrrolidin-1-ylcarbonyl, N-(methyl)-N-(dimethylaminocarbonylmethyl)-aminocarbonyl, (3-oxo-piperazin-1-yl)carbonyl, N-(methyl)-N-(hydroxyethyl)-aminocarbonyl, 2-(2-oxo-imidazolidin-1-yl)-

20 ethylaminocarbonyl, methylaminocarbonylmethylaminocarbonyl, tetrahydropyran-4-ylmethylaminocarbonyl, (4-hydroxypiperidin-1-yl)carbonyl, (4-hydroxyethyl-piperazin-1-yl)carbonyl, N-(1-methylpiperidin-4-yl)-N-(methyl)-aminocarbonyl, imidazol-1-ylpropylaminocarbonyl, 4-(pyrrolidin-1-yl)piperidin-1-ylcarbonyl, methoxyethylaminocarbonyl, cyclopropylmethyaminocarbonyl,

25 methylsulfonylethylaminocarbonyl, 2-(2-oxo-pyrrolidin-1-yl)-ethylaminocarbonyl, 3-hydroxylazetidin-1-ylcarbonyl, morpholinocarbonyl, N-acylpiperazin-1-ylcarbonyl, (N-methylpiperidin-4-yl)aminocarbonyl, imidazol-2-ylmethylaminocarbonyl, azetidin-1-ylcarbonyl, N-methylhomopiperazin-1-ylcarbonyl, dimethylaminocarbonyl, aminosulfonyl, dimethylaminosulfonyl, isopropylaminosulfonyl, N-methylpiperazin-1-ylsulfonyl,

30 methylsulfinyl, methylthio, 1,3,4-oxadiazoly-2-yl, 2,5-dimethylisoxazol-4-yl, 3-furyl, and methylsulfonyl;

R³ is chloro or fluoro;

or a salt, pro-drug or solvate thereof.

Further preferred compounds of the invention are each of the Examples, each of which provides a further independent aspect of the invention. In further aspects, the present invention also comprises any two or more compounds of the Examples.

5 In one aspect, particular compounds of the invention comprise any one or more of:
3-(1-methylethyl)oxy-5-{4-[(4-methylpiperazin-1-yl)carbonyl]phenoxy}-N-1,3-thiazol-2-
ylbenzamide;
1-(4-{3-(1-methylethyl)oxy-5-[(1,3-thiazol-2-ylamino)carbonyl]phenoxybenzoyl}prolinamide;
3-(4-{{[2-(dimethylamino)-2-oxoethyl](methyl)amino]carbonyl}phenoxy)-5-(1-
10 methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;
3-(1-methylethyl)oxy-5-{4-[(3-oxopiperazin-1-yl)carbonyl]phenoxy}-N-1,3-thiazol-2-
ylbenzamide;
3-(4-{{(2-hydroxyethyl)(methyl)amino]carbonyl}phenoxy)-5-(1-methylethyl)oxy-N-1,3-
thiazol-2-ylbenzamide;
15 3-(4-{{(2-hydroxyethyl)amino]carbonyl}phenoxy)-5-(1-methylethyl)oxy-N-1,3-thiazol-2-
ylbenzamide;
3-(1-methylethyl)oxy-5-[4-({{[2-(2-oxoimidazolidin-1-yl)ethyl]amino}carbonyl}phenoxy]-N-
1,3-thiazol-2-ylbenzamide;
3-(1-methylethyl)oxy-5-[4-({[2-(methylamino)-2-oxoethyl]amino}carbonyl)phenoxy]-N-1,3-
20 thiazol-2-ylbenzamide;
3-(1-methylethyl)oxy-5-(4-{{[(tetrahydro-2H-pyran-4-yl)methyl]amino}carbonyl}phenoxy)-N-
1,3-thiazol-2-ylbenzamide;
3-{4-[(4-hydroxypiperidin-1-yl)carbonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-
ylbenzamide;
25 3-(4-{{[4-(2-hydroxyethyl)piperazin-1-yl]carbonyl}phenoxy)-5-(1-methylethyl)oxy-N-1,3-
thiazol-2-ylbenzamide;
3-(1-methylethyl)oxy-5-(4-{{[methyl(1-methylpiperidin-4-yl)amino]carbonyl}phenoxy)-N-1,3-
thiazol-2-ylbenzamide;
3-[4-{{[3-(1H-imidazol-1-yl)propyl]amino}carbonyl}phenoxy]-5-(1-methylethyl)oxy-N-1,3-
30 thiazol-2-ylbenzamide;
3-(1-methylethyl)oxy-5-{4-[(4-pyrrolidin-1-yl)piperidin-1-yl]carbonyl}phenoxy)-N-1,3-
thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-(4-{[(2-methoxyethyl)amino]carbonyl}phenoxy)-N-1,3-thiazol-2-ylbenzamide;

3-(4-{[(cyclopropylmethyl)amino]carbonyl}phenoxy)-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

5 3-(1-methylethyl)oxy-5-[4-({[2-(methylsulfonyl)ethyl]amino}carbonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-({[2-(2-oxopyrrolidin-1-yl)ethyl]amino}carbonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-{4-[(3-hydroxyazetidin-1-yl)carbonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

10 3-(1-methylethyl)oxy-5-[4-(morpholin-4-ylcarbonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-{4-[(4-acetylpirperazin-1-yl)carbonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-(4-{[(1-methylpiperidin-4-yl)amino]carbonyl}phenoxy)-N-1,3-thiazol-2-ylbenzamide;

15 3-(4-{[(1H-imidazol-2-yl)methyl]amino}carbonyl)phenoxy)-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-{{[4-(azetidin-1-ylcarbonyl)phenyl]oxy}-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide};

20 3-chloro-4-{3-(1-methylethyl)oxy-5-[(1,3-thiazol-2-ylamino)carbonyl]phenoxy}-N-(2-methoxyethyl)benzamide;

3-chloro-4-{3-(1-methylethyl)oxy-5-[(1,3-thiazol-2-ylamino)carbonyl]phenoxy}-N,N-dimethylbenzamide;

3-[4-(aminosulfonyl)-2-fluorophenoxy]-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

25 3-{2-chloro-4-[(dimethylamino)sulfonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-{2-chloro-4-[((1-methylethyl)amino)sulfonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-{2-chloro-4-[(4-methylpirperazin-1-yl)sulfonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

30 3-[4-(aminosulfonyl)-5-chloro-2-fluorophenoxy]-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-(methylsulfinyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-[4-(ethylthio)phenoxy]-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-(1,3,4-oxadiazol-2-yl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-[4-(3,5-dimethylisoxazol-4-yl)phenoxy]-5-(1-methylethyl)oxy-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

5 3-[(4-furan-3-ylphenyl)oxy]-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

3-(1-methylethyl)oxy-N-(1-methyl-1H-pyrazol-3-yl)-5-[4-(methylsulfonyl)phenoxy]benzamide;

3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]-N-1,3,4-thiadiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

10 3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]-N-pyridin-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]-N-pyrazin-2-ylbenzamide;

3-(1-methylethyl)oxy-N-(5-methylisoxazol-3-yl)-5-[4-(methylsulfonyl)phenoxy]benzamide;

3-(1-methylethyl)oxy-N-isoxazol-3-yl-5-[4-(methylsulfonyl)phenoxy]benzamide;

N-[5-(2-furyl)-1,3,4-thiadiazol-2-yl]-3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]benzamide; and

15 N-{4-[(dimethylamino)methyl]-1,3-thiazol-2-yl}-3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]benzamide;

3-{{[4-(azetidin-1-ylcarbonyl)-2-chlorophenyl]oxy}-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

20 3-{{[4-(azetidin-1-ylcarbonyl)-2-fluorophenyl]oxy}-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

3-{{[4-(azetidin-1-ylcarbonyl)phenyl]oxy}-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

3-[(1-methylethyl)oxy]-5-[(4-{{[methyl(1-methylpiperidin-4-yl)amino]carbonyl}phenyl})oxy]-

25 N-(3-methyl-1,2,4-thiadiazol-5-yl)benzamide;

3-({4-[(4-methyl-1,4-diazepan-1-yl)carbonyl]phenyl})oxy)-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide;

or a salt, pro-drug or solvate thereof.

30 In another aspect, particular compounds of the invention comprise any one or more of:

3-(4-{{[[2-(dimethylamino)-2-oxoethyl](methyl)amino]carbonyl}phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(4-{[(2-hydroxyethyl)(methyl)amino]carbonyl}phenoxy)-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(4-{[(2-hydroxyethyl)amino]carbonyl}phenoxy)-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

5 3-(1-methylethyl)oxy-5-[4-({[2-(2-oxoimidazolidin-1-yl)ethyl]amino}carbonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-({[2-(methylamino)-2-oxoethyl]amino}carbonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-(4-{{[(tetrahydro-2H-pyran-4-yl)methyl]amino}carbonyl}phenoxy)-N-10 1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-(4-{{[methyl(1-methylpiperidin-4-yl)amino]carbonyl}phenoxy}-N-1,3-thiazol-2-ylbenzamide;

3-[4-({[3-(1H-imidazol-1-yl)propyl]amino}carbonyl)phenoxy]-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

15 3-(1-methylethyl)oxy-5-(4-{{[(2-methoxyethyl)amino]carbonyl}phenoxy}-N-1,3-thiazol-2-ylbenzamide;

3-(4-{{[(cyclopropylmethyl)amino]carbonyl}phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-({[2-(methylsulfonyl)ethyl]amino}carbonyl)phenoxy]-N-1,3-20 thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-({[2-(2-oxopyrrolidin-1-yl)ethyl]amino}carbonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-(4-{{[(1-methylpiperidin-4-yl)amino]carbonyl}phenoxy}-N-1,3-thiazol-2-ylbenzamide;

25 3-(4-{{[(1H-imidazol-2-yl)methyl]amino}carbonyl}phenoxy)-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-chloro-4-{3-(1-methylethyl)oxy-5-[(1,3-thiazol-2-ylamino)carbonyl]phenoxy}-N-(2-methoxyethyl)benzamide;

3-[(1-methylethyl)oxy]-5-[(4-{{[methyl(1-methylpiperidin-4-yl)amino]carbonyl}phenyl}oxy)-30 N-(3-methyl-1,2,4-thiadiazol-5-yl)benzamide;

or a salt, pro-drug or solvate thereof.

In another aspect, particular compounds of the invention comprise any one or more of:

3-(1-methylethyl)oxy-5-{4-[(4-methylpiperazin-1-yl)carbonyl]phenoxy}-N-1,3-thiazol-2-ylbenzamide;

1-(4-{3-(1-methylethyl)oxy-5-[(1,3-thiazol-2-ylamino)carbonyl]phenoxy}benzoyl)prolinamide;

5 3-(1-methylethyl)oxy-5-{4-[(3-oxopiperazin-1-yl)carbonyl]phenoxy}-N-1,3-thiazol-2-ylbenzamide;

3-{4-[(4-hydroxypiperidin-1-yl)carbonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(4-{[4-(2-hydroxyethyl)piperazin-1-yl]carbonyl}phenoxy)-5-(1-methylethyl)oxy-N-1,3-

10 thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-{4-[(4-pyrrolidin-1-yl)piperidin-1-yl]carbonyl}phenoxy}-N-1,3-thiazol-2-ylbenzamide;

3-{4-[(3-hydroxyazetidin-1-yl)carbonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

15 3-(1-methylethyl)oxy-5-[4-(morpholin-4-ylcarbonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-{4-[(4-acetyl)piperazin-1-yl)carbonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-{[4-(azetidin-1-ylcarbonyl)phenyl]oxy}-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide;

20 3-{[4-[(4-methyl-1,4-diazepan-1-yl)carbonyl]phenyl]oxy}-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide;

3-{2-chloro-4-[(4-methylpiperazin-1-yl)sulfonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-{[4-(azetidin-1-ylcarbonyl)-2-chlorophenyl]oxy}-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-

25 pyrazol-3-yl)benzamide;

3-{[4-(azetidin-1-ylcarbonyl)-2-fluorophenyl]oxy}-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

3-{[4-(azetidin-1-ylcarbonyl)phenyl]oxy}-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

30 or a salt, pro-drug or solvate thereof.

In another aspect, particular compounds of the invention comprise any one or more of:

3-(1-methylethyl)oxy-5-{4-[(4-methylpiperazin-1-yl)carbonyl]phenoxy}-N-1,3-thiazol-2-ylbenzamide;

3-{4-[(4-hydroxypiperidin-1-yl)carbonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(4-[(4-(2-hydroxyethyl)piperazin-1-yl)carbonyl]phenoxy)-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

5 3-[(4-[(4-methyl-1,4-diazepan-1-yl)carbonyl]phenyl)oxy]-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide;

3-[(4-(azetidin-1-ylcarbonyl)-2-chlorophenyl)oxy]-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

3-[(4-(azetidin-1-ylcarbonyl)-2-fluorophenyl)oxy]-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

10 or a salt, pro-drug or solvate thereof.

15 In another aspect, particular compounds of the invention comprise any one or more of:

3-(1-methylethyl)oxy-5-[4-(1,3,4-oxadiazol-2-yl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-[4-(3,5-dimethylisoxazol-4-yl)phenoxy]-5-(1-methylethyl)oxy-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

3-[(4-furan-3-ylphenyl)oxy]-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide;

20 or a salt, pro-drug or solvate thereof.

In another aspect, particular compounds of the invention comprises:

N-[5-(2-furyl)-1,3,4-thiadiazol-2-yl]-3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]benzamide

25 or a salt, pro-drug or solvate thereof.

In another aspect, particular compounds of the invention comprise any one or more of:

3-{4-[(dimethylamino)carbonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

30 3-(1-methylethyl)oxy-5-{4-[(methylamino)carbonyl]phenoxy}-N-1,3-thiazol-2-ylbenzamide;

or a salt, pro-drug or solvate thereof.

In another aspect, particular compounds of the invention comprise any one or more of:

3-chloro-4-{3-(1-methylethyl)oxy-5-[(1,3-thiazol-2-ylamino)carbonyl]phenoxy}-N,N-dimethylbenzamide;

3-[4-(aminosulfonyl)-2-fluorophenoxy]-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-{2-chloro-4-[(dimethylamino)sulfonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

5 3-{2-chloro-4-[((1-methylethyl)amino)sulfonyl]phenoxy}-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-[4-(aminosulfonyl)-5-chloro-2-fluorophenoxy]-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

10 3-(1-methylethyl)oxy-5-[4-(methylsulfinyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-[4-(ethylthio)phenoxy]-5-(1-methylethyl)oxy-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-N-(1-methyl-1H-pyrazol-3-yl)-5-[4-(methylsulfonyl)phenoxy]benzamide;

3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]-N-1,3,4-thiadiazol-2-ylbenzamide;

15 3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]-N-1,3-thiazol-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]-N-pyridin-2-ylbenzamide;

3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]-N-pyrazin-2-ylbenzamide;

3-(1-methylethyl)oxy-N-(5-methylisoxazol-3-yl)-5-[4-(methylsulfonyl)phenoxy]benzamide;

3-(1-methylethyl)oxy-N-isoxazol-3-yl-5-[4-(methylsulfonyl)phenoxy]benzamide;

20 N-{4-[(dimethylamino)methyl]-1,3-thiazol-2-yl}-3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]benzamide;

or a salt, pro-drug or solvate thereof.

The compounds of the invention may be administered in the form of a pro-drug. A pro-drug is a bioprecursor or pharmaceutically acceptable compound being degradable in the body to produce a compound of the invention (such as an ester or amide of a compound of the invention, particularly an in-vivo hydrolysable ester). Various forms of prodrugs are known in the art. For examples of such prodrug derivatives, see:

a) Design of Prodrugs, edited by H. Bundgaard, (Elsevier, 1985) and Methods in Enzymology, Vol. 42, p. 309-396, edited by K. Widder, *et al.* (Academic Press, 1985);

b) A Textbook of Drug Design and Development, edited by Krogsgaard-Larsen;

c) H. Bundgaard, Chapter 5 "Design and Application of Prodrugs", by H. Bundgaard p. 113-191 (1991);

- d) H. Bundgaard, Advanced Drug Delivery Reviews, 8, 1-38 (1992);
- e) H. Bundgaard, *et al.*, Journal of Pharmaceutical Sciences, 77, 285 (1988); and
- f) N. Kakeya, *et al.*, Chem Pharm Bull, 32, 692 (1984).

The contents of the above cited documents are incorporated herein by reference.

5 Examples of pro-drugs are as follows. An in-vivo hydrolysable ester of a compound of the invention containing a carboxy or a hydroxy group is, for example, a pharmaceutically-acceptable ester which is hydrolysed in the human or animal body to produce the parent acid or alcohol. Suitable pharmaceutically-acceptable esters for carboxy include C₁ to C₆alkoxymethyl esters for example methoxymethyl, C₁ to C₆alkanoyloxymethyl esters 10 for example pivaloyloxymethyl, phthalidyl esters, C₃ to C₈cycloalkoxycarbonyloxyC₁ to C₆alkyl esters for example 1-cyclohexylcarbonyloxyethyl; 1,3-dioxolen-2-onylmethyl esters, for example 5-methyl-1,3-dioxolen-2-onylmethyl; and C₁-alkoxycarbonyloxyethyl esters.

An in-vivo hydrolysable ester of a compound of the invention containing a hydroxy 15 group includes inorganic esters such as phosphate esters (including phosphoramidic cyclic esters) and α-acyloxyalkyl ethers and related compounds which as a result of the in-vivo hydrolysis of the ester breakdown to give the parent hydroxy group/s. Examples of α-acyloxyalkyl ethers include acetoxymethoxy and 2,2-dimethylpropionyloxy-methoxy. A selection of in-vivo hydrolysable ester forming groups for hydroxy include alkanoyl, 20 benzoyl, phenylacetyl and substituted benzoyl and phenylacetyl, alkoxy carbonyl (to give alkyl carbonate esters), dialkylcarbamoyl and N-(dialkylaminoethyl)-N-alkylcarbamoyl (to give carbamates), dialkylaminoacetyl and carboxyacetyl.

A suitable pharmaceutically-acceptable salt of a compound of the invention is, for example, an acid-addition salt of a compound of the invention which is sufficiently basic, for 25 example, an acid-addition salt with, for example, an inorganic or organic acid, for example hydrochloric, hydrobromic, sulphuric, phosphoric, trifluoroacetic, citric or maleic acid. In addition a suitable pharmaceutically-acceptable salt of a benzoxazinone derivative of the invention which is sufficiently acidic is an alkali metal salt, for example a sodium or potassium salt, an alkaline earth metal salt, for example a calcium or magnesium salt, an 30 ammonium salt or a salt with an organic base which affords a physiologically-acceptable cation, for example a salt with methylamine, dimethylamine, trimethylamine, piperidine, morpholine or tris-(2-hydroxyethyl)amine.

A further feature of the invention is a pharmaceutical composition comprising a compound of Formula (I) as defined above, or a salt, solvate or prodrug thereof, together with a pharmaceutically-acceptable diluent or carrier.

According to another aspect of the invention there is provided a compound of Formula 5 (I) as defined above for use as a medicament.

Further according to the invention there is provided a compound of Formula (I) for use in the preparation of a medicament for treatment of a disease mediated through GLK, in particular type 2 diabetes.

The compound is suitably formulated as a pharmaceutical composition for use in this 10 way.

According to another aspect of the present invention there is provided a method of treating GLK mediated diseases, especially diabetes, by administering an effective amount of a compound of Formula (I) or salt, solvate or pro-drug thereof, to a mammal in need of such treatment.

15 Specific diseases which may be treated by a compound or composition of the invention include: blood glucose lowering in type 2 Diabetes Mellitus without a serious risk of hypoglycaemia (and potential to treat type 1), dyslipidemia, obesity, insulin resistance, metabolic syndrome X, impaired glucose tolerance.

As discussed above, thus the GLK/GLKRP system can be described as a potential 20 "Diabesity" target (of benefit in both Diabetes and Obesity). Thus, according to another aspect of the invention there is provided the use of a compound of Formula (I) or salt, solvate or pro-drug thereof, in the preparation of a medicament for use in the combined treatment or prevention of diabetes and obesity.

According to another aspect of the invention there is provided the use of a compound of 25 Formula (I) or salt, solvate or pro-drug thereof, in the preparation of a medicament for use in the treatment or prevention of obesity.

According to a further aspect of the invention there is provided a method for the combined treatment of obesity and diabetes by administering an effective amount of a compound of Formula (I) or salt, solvate or pro-drug thereof, to a mammal in need of such 30 treatment.

According to a further aspect of the invention there is provided a method for the treatment of obesity by administering an effective amount of a compound of Formula (I) or salt, solvate or pro-drug thereof, to a mammal in need of such treatment.

The compositions of the invention may be in a form suitable for oral use (for example as tablets, lozenges, hard or soft capsules, aqueous or oily suspensions, emulsions, dispersible powders or granules, syrups or elixirs), for topical use (for example as creams, ointments, gels, or aqueous or oily solutions or suspensions), for administration by inhalation (for example as a finely divided powder or a liquid aerosol), for administration by insufflation (for example as a finely divided powder) or for parenteral administration (for example as a sterile aqueous or oily solution for intravenous, subcutaneous, intramuscular or intramuscular dosing or as a suppository for rectal dosing). Dosage forms suitable for oral use are preferred.

The compositions of the invention may be obtained by conventional procedures using conventional pharmaceutical excipients, well known in the art. Thus, compositions intended for oral use may contain, for example, one or more colouring, sweetening, flavouring and/or preservative agents.

Suitable pharmaceutically acceptable excipients for a tablet formulation include, for example, inert diluents such as lactose, sodium carbonate, calcium phosphate or calcium carbonate, granulating and disintegrating agents such as corn starch or algenic acid; binding agents such as starch; lubricating agents such as magnesium stearate, stearic acid or talc; preservative agents such as ethyl or propyl p-hydroxybenzoate, and anti-oxidants, such as ascorbic acid. Tablet formulations may be uncoated or coated either to modify their disintegration and the subsequent absorption of the active ingredient within the gastrointestinal tract, or to improve their stability and/or appearance, in either case, using conventional coating agents and procedures well known in the art.

Compositions for oral use may be in the form of hard gelatin capsules in which the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules in which the active ingredient is mixed with water or an oil such as peanut oil, liquid paraffin, or olive oil.

Aqueous suspensions generally contain the active ingredient in finely powdered form together with one or more suspending agents, such as sodium carboxymethylcellulose, methylcellulose, hydroxypropylmethylcellulose, sodium alginate, polyvinyl-pyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents such as lecithin or condensation products of an alkylene oxide with fatty acids (for example polyoxethylene stearate), or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or

condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and

5 hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives (such as ethyl or propyl p-hydroxybenzoate, anti-oxidants (such as ascorbic acid), colouring agents, flavouring agents, and/or sweetening agents (such as sucrose, saccharine or aspartame).

Oily suspensions may be formulated by suspending the active ingredient in a vegetable oil (such as arachis oil, olive oil, sesame oil or coconut oil) or in a mineral oil (such as liquid paraffin). The oily suspensions may also contain a thickening agent such as beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set out above, and flavouring agents may be added to provide a palatable oral preparation. These compositions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

15 Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water generally contain the active ingredient together with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above. Additional excipients such as sweetening, flavouring and colouring agents, may also be present.

20 The pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, such as olive oil or arachis oil, or a mineral oil, such as for example liquid paraffin or a mixture of any of these. Suitable emulsifying agents may be, for example, naturally-occurring gums such as gum acacia or gum tragacanth, naturally-occurring phosphatides such as soya bean, lecithin, an esters or partial 25 esters derived from fatty acids and hexitol anhydrides (for example sorbitan monooleate) and condensation products of the said partial esters with ethylene oxide such as polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening, flavouring and preservative agents.

Syrups and elixirs may be formulated with sweetening agents such as glycerol, 30 propylene glycol, sorbitol, aspartame or sucrose, and may also contain a demulcent, preservative, flavouring and/or colouring agent.

The pharmaceutical compositions may also be in the form of a sterile injectable aqueous or oily suspension, which may be formulated according to known procedures using one or

more of the appropriate dispersing or wetting agents and suspending agents, which have been mentioned above. A sterile injectable preparation may also be a sterile injectable solution or suspension in a non-toxic parenterally-acceptable diluent or solvent, for example a solution in 1,3-butanediol.

5 Compositions for administration by inhalation may be in the form of a conventional pressurised aerosol arranged to dispense the active ingredient either as an aerosol containing finely divided solid or liquid droplets. Conventional aerosol propellants such as volatile fluorinated hydrocarbons or hydrocarbons may be used and the aerosol device is conveniently arranged to dispense a metered quantity of active ingredient.

10 For further information on formulation the reader is referred to Chapter 25.2 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The amount of active ingredient that is combined with one or more excipients to produce a single dosage form will necessarily vary depending upon the host treated and the 15 particular route of administration. For example, a formulation intended for oral administration to humans will generally contain, for example, from 0.5 mg to 2 g of active agent compounded with an appropriate and convenient amount of excipients which may vary from about 5 to about 98 percent by weight of the total composition. Dosage unit forms will generally contain about 1 mg to about 500 mg of an active ingredient. For further information 20 on Routes of Administration and Dosage Regimes the reader is referred to Chapter 25.3 in Volume 5 of Comprehensive Medicinal Chemistry (Corwin Hansch; Chairman of Editorial Board), Pergamon Press 1990.

The size of the dose for therapeutic or prophylactic purposes of a compound of the Formula (I) will naturally vary according to the nature and severity of the conditions, the age 25 and sex of the animal or patient and the route of administration, according to well known principles of medicine.

In using a compound of the Formula (I) for therapeutic or prophylactic purposes it will generally be administered so that a daily dose in the range, for example, 0.5 mg to 75 mg per kg body weight is received, given if required in divided doses. In general lower doses will be 30 administered when a parenteral route is employed. Thus, for example, for intravenous administration, a dose in the range, for example, 0.5 mg to 30 mg per kg body weight will generally be used. Similarly, for administration by inhalation, a dose in the range, for

example, 0.5 mg to 25 mg per kg body weight will be used. Oral administration is however preferred.

The elevation of GLK activity described herein may be applied as a sole therapy or in combination with one or more other substances and/or treatments for the indication being 5 treated. Such conjoint treatment may be achieved by way of the simultaneous, sequential or separate administration of the individual components of the treatment. Simultaneous treatment may be in a single tablet or in separate tablets. For example in the treatment of diabetes mellitus, chemotherapy may include the following main categories of treatment:

- 1) Insulin and insulin analogues;
- 10 2) Insulin secretagogues including sulphonylureas (for example glibenclamide, glipizide), prandial glucose regulators (for example repaglinide, nateglinide);
- 3) Agents that improve incretin action (for example dipeptidyl peptidase IV inhibitors, and GLP-1 agonists);
- 4) Insulin sensitising agents including PPARgamma agonists (for example pioglitazone 15 and rosiglitazone), and agents with combined PPARalpha and gamma activity;
- 5) Agents that modulate hepatic glucose balance (for example metformin, fructose 1, 6 bisphosphatase inhibitors, glycogen phosphorylase inhibitors, glycogen synthase kinase inhibitors);
- 6) Agents designed to reduce the absorption of glucose from the intestine (for example 20 acarbose);
- 7) Agents that prevent the reabsorption of glucose by the kidney (SGLT inhibitors);
- 8) Agents designed to treat the complications of prolonged hyperglycaemia (for example aldose reductase inhibitors);
- 9) Anti-obesity agents (for example sibutramine and orlistat);
- 25 10) Anti-dyslipidaemia agents such as, HMG-CoA reductase inhibitors (eg statins); PPAR α agonists (fibrates, eg gemfibrozil); bile acid sequestrants (cholestyramine); cholesterol absorption inhibitors (plant stanols, synthetic inhibitors); bile acid absorption inhibitors (IBATi) and nicotinic acid and analogues (niacin and slow release formulations);
- 30 11) Antihypertensive agents such as, β blockers (eg atenolol, inderal); ACE inhibitors (eg lisinopril); Calcium antagonists (eg. nifedipine); Angiotensin receptor antagonists (eg candesartan), α antagonists and diuretic agents (eg. furosemide, benzthiazide);

12) Haemostasis modulators such as, antithrombotics, activators of fibrinolysis and antiplatelet agents; thrombin antagonists; factor Xa inhibitors; factor VIIa inhibitors); antiplatelet agents (eg. aspirin, clopidogrel); anticoagulants (heparin and Low molecular weight analogues, hirudin) and warfarin;

5 13) Agents which antagonise the actions of glucagon; and

14) Anti-inflammatory agents, such as non-steroidal anti-inflammatory drugs (eg. aspirin) and steroidal anti-inflammatory agents (eg. cortisone).

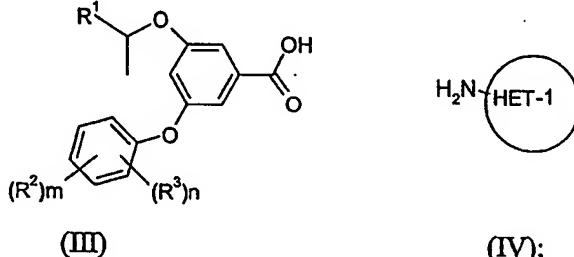
According to another aspect of the present invention there is provided individual compounds produced as end products in the Examples set out below and salts, solvates and
10 pro-drugs thereof.

A compound of the invention, or a salt thereof, may be prepared by any process known to be applicable to the preparation of such compounds or structurally related compounds. Functional groups may be protected and deprotected using conventional methods. For examples of protecting groups such as amino and carboxylic acid protecting groups (as well as
15 means of formation and eventual deprotection), see T.W. Greene and P.G.M. Wuts, "Protective Groups in Organic Synthesis", Second Edition, John Wiley & Sons, New York, 1991.

Processes for the synthesis of compounds of Formula (I) are provided as a further feature of the invention. Thus, according to a further aspect of the invention there is provided
20 a process for the preparation of a compound of Formula (I), which comprises a process a) to d) (wherein the variables are as defined hereinbefore for compounds of Formula (I) unless otherwise defined):

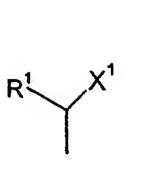
(a) reaction of an acid of Formula (III) or activated derivative thereof with a compound of Formula (IV),

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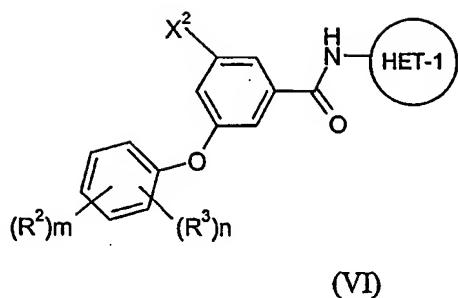


or

(b) reaction of a compound of Formula (V) with a compound of Formula (VI),



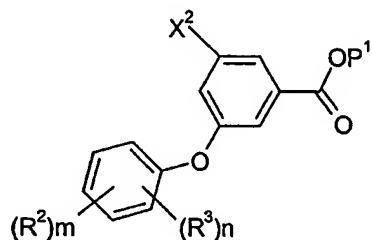
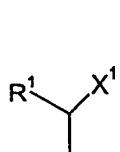
(V)



(VI)

wherein X^1 is a leaving group and X^2 is a hydroxyl group or X^1 is a hydroxyl group and X^2 is a leaving group;

5 process (b) could also be accomplished using the intermediate ester Formula (VII), wherein P^1 is a protecting group as hereinafter described, followed by ester hydrolysis and amide formation by procedures described elsewhere and well known to those skilled in the art;

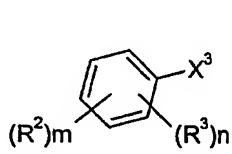


(VII)

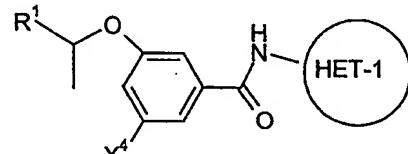
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or

(c) reaction of a compound of Formula (VIII) with a compound of Formula (IX)

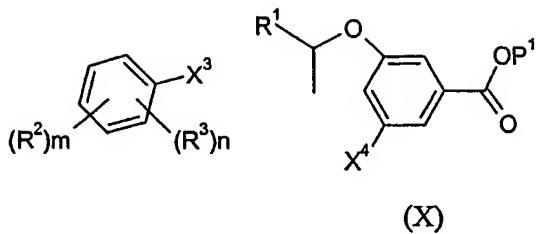


(VIII)



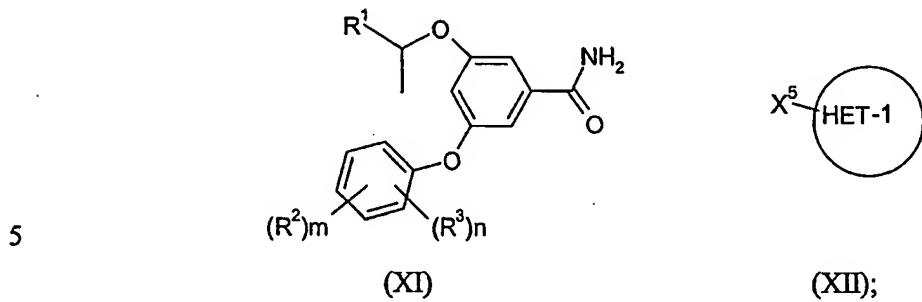
(IX)

15 wherein X^3 is a leaving group or an organometallic reagent and X^4 is a hydroxyl group or X^3 is a hydroxyl group and X^4 is a leaving group or an organometallic reagent; process (c) could also be accomplished using the intermediate ester Formula (X), followed by ester hydrolysis and amide formation by procedures described elsewhere and well known to those skilled in the art;



or

(d) reaction of a compound of Formula (XI) with a compound of Formula (XII),



wherein X⁵ is a leaving group;

and thereafter, if necessary:

10 i) converting a compound of Formula (I) into another compound of Formula (I);
ii) removing any protecting groups; and/or
iii) forming a salt, pro-drug or solvate thereof.

Suitable leaving groups X¹ to X⁵ for processes b) to d) are any leaving group known in the art for these types of reactions, for example halo, alkoxy, trifluoromethanesulfonyloxy, methanesulfonyloxy, or p-toluenesulfonyloxy, or a group (such as a hydroxy group) that could be converted into a leaving group (such as an oxytriphenylphosphonium group) *in situ*.

Compounds of Formulae (III) to (XII) are commercially available, or are known in the art, or may be made by processes known in the art as shown, for example, in the accompanying Examples. For further information on processes for making such compounds, we refer to our PCT publications WO 03/000267, WO 03/015774 and WO 03/000262 and references therein. In general it will be appreciated that any aryl-O or alkyl-O bond may be formed by nucleophilic substitution or metal catalysed processes, optionally in the presence of a suitable base.

Examples of conversions of a compound of Formula (I) into another compound of Formula (I), well known to those skilled in the art, include functional group interconversions such as hydrolysis, hydrogenation, hydrogenolysis, oxidation or reduction, and/or further 25 functionalisation by standard reactions such as amide or metal-catalysed coupling, or nucleophilic displacement reactions:

Specific reaction conditions for the above reactions are as follows, wherein when P¹ is a protecting group P¹ is preferably C₁₋₄alkyl, for example methyl or ethyl:

Process a) – coupling reactions of amino groups with carboxylic acids to form an amide are well known in the art. For example,

- 5 (i) using an appropriate coupling reaction, such as a carbodiimide coupling reaction performed with EDAC in the presence of DMAP in a suitable solvent such as DCM, chloroform or DMF at room temperature; or
(ii) reaction in which the carboxylic group is activated to an acid chloride by reaction with oxalyl chloride in the presence of a suitable solvent such as methylene chloride. The acid
- 10 chloride can then be reacted with a compound of Formula (IV) in the presence of a base, such as triethylamine or pyridine, in a suitable solvent such as chloroform or DCM at a temperature between 0°C and 80°C.

Process b) – compounds of Formula (V) and (VI) can be reacted together in a suitable solvent, such as DMF or THF, with a base such as sodium hydride or potassium *tert*-butoxide, at a

- 15 temperature in the range 0 to 100°C, optionally using microwave heating or metal catalysis such as palladium(II)acetate, palladium on carbon, copper(II)acetate or copper(I)iodide; Alternatively, compounds of Formula (V) and (VI) can be reacted together in a suitable solvent, such as THF or DCM, with a suitable phosphine such as triphenylphosphine, and azodicarboxylate such as diethylazodicarboxylate; process b) could also be carried out using a
- 20 precursor to the ester of formula (VII) such as an aryl-nitrile or trifluoromethyl derivative, followed by conversion to a carboxylic acid and amide formation as previously described;

Process c) – compounds of Formula (VIII) and (IX) can be reacted together in a suitable solvent, such as DMF or THF, with a base such as sodium hydride or potassium *tert*-butoxide, at a temperature in the range 0 to 200°C, optionally using microwave heating or metal

 - 25 catalysis such as palladium(II)acetate, palladium on carbon, copper(II)acetate or copper(I)iodide; process c) could also be carried out using a precursor to the ester of formula (X) such as an aryl-nitrile or trifluoromethyl derivative, followed by conversion to a carboxylic acid and amide formation as previously described;

Process d) – reaction of a compound of Formula (XI) with a compound of Formula (XII) can

 - 30 be performed in a polar solvent, such as DMF or a non-polar solvent such as THF with a strong base, such as sodium hydride or potassium *tert*-butoxide at a temperature between 0

and 200°C, optionally using microwave heating or metal catalysis, such as palladium(II)acetate, palladium on carbon, copper(II)acetate or copper(I)iodide.

Certain intermediates of formula (III), (VI), (VII), (IX) and/or (XI) are believed to be novel and comprise an independent aspect of the invention.

5 During the preparation process, it may be advantageous to use a protecting group for a functional group within the molecule. Protecting groups may be removed by any convenient method as described in the literature or known to the skilled chemist as appropriate for the removal of the protecting group in question, such methods being chosen so as to effect removal of the protecting group with minimum disturbance of groups elsewhere in the
10 molecule.

Specific examples of protecting groups are given below for the sake of convenience, in which "lower" signifies that the group to which it is applied preferably has 1-4 carbon atoms. It will be understood that these examples are not exhaustive. Where specific examples of methods for the removal of protecting groups are given below these are similarly not
15 exhaustive. The use of protecting groups and methods of deprotection not specifically mentioned is of course within the scope of the invention.

A carboxy protecting group may be the residue of an ester-forming aliphatic or araliphatic alcohol or of an ester-forming silanol (the said alcohol or silanol preferably containing 1-20 carbon atoms). Examples of carboxy protecting groups include straight or
20 branched chain (1-12C)alkyl groups (e.g. isopropyl, *t*-butyl); lower alkoxy lower alkyl groups (e.g. methoxymethyl, ethoxymethyl, isobutoxymethyl; lower aliphatic acyloxy lower alkyl groups, (e.g. acetoxymethyl, propionyloxymethyl, butyryloxymethyl, pivaloyloxymethyl); lower alkoxycarbonyloxy lower alkyl groups (e.g. 1-methoxycarbonyloxyethyl, 1-ethoxycarbonyloxyethyl); aryl lower alkyl groups (e.g. *p*-methoxybenzyl, *o*-nitrobenzyl,
25 *p*-nitrobenzyl, benzhydryl and phthalidyl); tri(lower alkyl)silyl groups (e.g. trimethylsilyl and *t*-butyldimethylsilyl); tri(lower alkyl)silyl lower alkyl groups (e.g. trimethylsilylethyl); and (2-6C)alkenyl groups (e.g. allyl and vinylthethyl).

Methods particularly appropriate for the removal of carboxyl protecting groups include for example acid-, metal- or enzymically-catalysed hydrolysis.
30 Examples of hydroxy protecting groups include lower alkenyl groups (e.g. allyl); lower alkanoyl groups (e.g. acetyl); lower alkoxycarbonyl groups (e.g. *t*-butoxycarbonyl); lower alkenyloxycarbonyl groups (e.g. allyloxycarbonyl); aryl lower alkoxycarbonyl groups (e.g. benzoyloxycarbonyl, *p*-methoxybenzyloxycarbonyl, *o*-nitrobenzyloxycarbonyl,

p-nitrobenzyloxycarbonyl); tri lower alkyl/arylsilyl groups (e.g. trimethylsilyl, t-butyldimethylsilyl, t-butyldiphenylsilyl); aryl lower alkyl groups (e.g. benzyl) groups; and triaryl lower alkyl groups (e.g. triphenylmethyl).

Examples of amino protecting groups include formyl, aralkyl groups (e.g. benzyl and

5 substituted benzyl, e.g. p-methoxybenzyl, nitrobenzyl and 2,4-dimethoxybenzyl, and triphenylmethyl); di-p-anisylmethyl and furylmethyl groups; lower alkoxy carbonyl (e.g. t-butoxycarbonyl); lower alkenyloxycarbonyl (e.g. allyloxycarbonyl); aryl lower alkoxy carbonyl groups (e.g. benzyloxycarbonyl, p-methoxybenzyloxycarbonyl,

10 o-nitrobenzyloxycarbonyl, p-nitrobenzyloxycarbonyl; trialkylsilyl (e.g. trimethylsilyl and t-butyldimethylsilyl); alkylidene (e.g. methylidene); benzylidene and substituted benzylidene groups.

Methods appropriate for removal of hydroxy and amino protecting groups include, for example, acid-, base, metal- or enzymically-catalysed hydrolysis, or photolytically for groups such as o-nitrobenzyloxycarbonyl, or with fluoride ions for silyl groups.

15 Examples of protecting groups for amide groups include aralkoxymethyl (e.g. benzyloxymethyl and substituted benzyloxymethyl); alkoxy methyl (e.g. methoxymethyl and trimethylsilylethoxymethyl); tri alkyl/arylsilyl (e.g. trimethylsilyl, t-butyldimethylsilyl, t-butyldiphenylsilyl); tri alkyl/arylsilyloxy methyl (e.g. t-butyldimethylsilyloxy methyl, t-butyldiphenylsilyloxy methyl); 4-alkoxyphenyl (e.g. 4-methoxyphenyl); 2,4-di(alkoxy)phenyl

20 (e.g. 2,4-dimethoxyphenyl); 4-alkoxybenzyl (e.g. 4-methoxybenzyl); 2,4-di(alkoxy)benzyl (e.g. 2,4-di(methoxy)benzyl); and alk-1-enyl (e.g. allyl, but-1-enyl and substituted vinyl e.g. 2-phenylvinyl).

Aralkoxymethyl groups may be introduced onto the amide group by reacting the latter group with the appropriate aralkoxymethyl chloride, and removed by catalytic hydrogenation.

25 Alkoxy methyl, tri alkyl/arylsilyl and tri alkyl/silyloxy methyl groups may be introduced by reacting the amide with the appropriate chloride and removing with acid; or in the case of the silyl containing groups, fluoride ions. The alkoxyphenyl and alkoxybenzyl groups are conveniently introduced by arylation or alkylation with an appropriate halide and removed by oxidation with ceric ammonium nitrate. Finally alk-1-enyl groups may be introduced by

30 reacting the amide with the appropriate aldehyde and removed with acid.

The following examples are for illustration purposes and are not intended to limit the scope of this application. Each exemplified compound represents a particular and

independent aspect of the invention. In the following non-limiting Examples, unless otherwise stated:

- (i) evaporation were carried out by rotary evaporation in *vacuo* and work-up procedures were carried out after removal of residual solids such as drying agents by filtration;
- 5 (ii) operations were carried out at room temperature, that is in the range 18-25°C and under an atmosphere of an inert gas such as argon or nitrogen;
- (iii) yields are given for illustration only and are not necessarily the maximum attainable;
- (iv) the structures of the end-products of the Formula (I) were confirmed by nuclear 10 (generally proton) magnetic resonance (NMR) with a field strength (for proton) of 300 or 400 MHz and mass spectral techniques; proton magnetic resonance chemical shift values were measured on the delta scale and peak multiplicities are shown as follows: s, singlet; d, doublet; t, triplet; m, multiplet; br, broad; q, quartet, quin, quintet;
- (v) intermediates were not generally fully characterised and purity was assessed by 15 thin layer chromatography (TLC), high-performance liquid chromatography (HPLC), infra-red (IR) or NMR analysis; and
- (vi) Biotage cartridges refer to pre-packed silica cartridges (from 40g up to 400g), eluted using a biotage pump and fraction collector system; Biotage UK Ltd, Hertford, Herts, UK.

20

Abbreviations

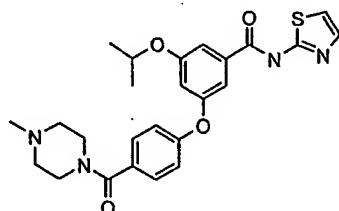
DCM	dichloromethane;
DEAD	diethylazodicarboxylate;
DIAD	diisopropylazodicarboxylate;
25 DMSO	dimethyl sulphoxide;
DMF	dimethylformamide;
EDAC	1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride;
HATU	<i>O</i> -(7-azabenzotriazol-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate;
30 HPLC	high pressure liquid chromatography;
HPMC	Hydroxypropylmethylcellulose;

LCMS	liquid chromatography / mass spectroscopy;
NMR	nuclear magnetic resonance spectroscopy;
RT	room temperature;
THF	tetrahydrofuran;
5 TFA	trifluoroacetic acid;
CDCl ₃	deuterochloroform

All compound names were derived using ACD NAME computer package.

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Example 1: 3-(1-Methylethyl)oxy-5-{4-[(4-methylpiperazin-1-yl)carbonyl]phenoxy}-N-1,3-thiazol-2-ylbenzamide

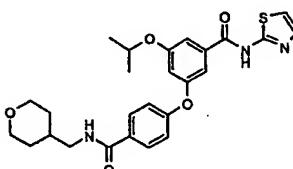
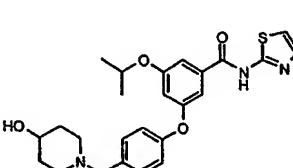
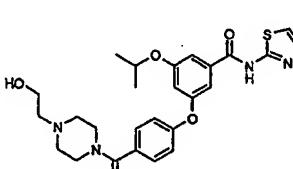
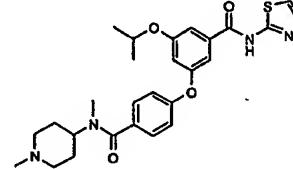
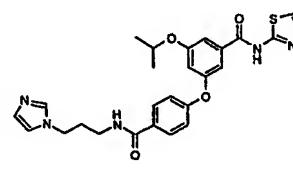
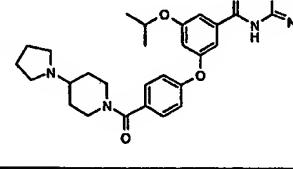
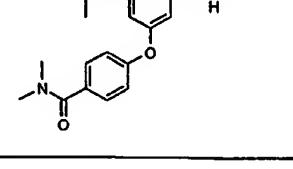


To a suspension of 4-((3-[(1-methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenyl)oxy)benzoic acid (100 mg), HATU (122 mg) and 1-methylpiperazine (32 mg) in DMF (2 mL) was added diisopropylethylamine (0.11 mL) and the mixture stirred at ambient temperature for 1 hour. Water (30 mL) was added and the mixture extracted with ethyl acetate. The combined organic extracts were washed with brine, dried (MgSO₄), and evaporated to a residue which was chromatographed on silica with 0-10% methanol in ethyl acetate as eluant 20 to give the desired compound (103 mg).

¹H NMR δ (d₆-DMSO): 1.3 (d, 6H), 2.15 (s, 3H), 2.3 (s, 4H), 3.4-3.5 (br, 4H), 4.7-4.8 (m, 1H), 6.85 (s, 1H), 7.1 (d, 2H), 7.25 (d, 1H), 7.3 (s, 1H), 7.4 (d, 2H), 7.5 (s, 1H), 7.55 (d, 1H); m/z 481 (M+H)⁺

25 In a similar manner, Examples 1a-1z were also prepared:

Example	Structure	<i>m/z</i>	NMR
1a		495 ($M+H$) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.35 (d, 6H), 1.8-1.95 (br, 3H), 2.2-2.25 (br, 1H), 3.5-3.7 (br, 2H), 4.4 (br, 1H), 4.8 (m, 1H), 6.9 (s, 1H), 6.95 (s, 1H), 7.1-7.2 (d, 2H), 7.35 (d, 1H), 7.4 (s, 1H), 7.55 (s, 1H), 7.6 (d, 1H), 7.7 (d, 1H), 12.63 (s, 1H)
1b		497 ($M+H$) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 2.65-3.0 (m, 9H), 4.1-4.3 (br, 2H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1-7.2 (d, 2H), 7.25 (d, 1H), 7.3 (m, 1H), 7.4-7.55 (m, 2H), 7.7 (d, 1H), 7.95 (s, 1H), 12.6 (s, 1H)
1c		481 ($M+H$) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 3.25 (m, 2H), 3.6-3.7 (br, 2H), 4.0 (m, 2H), 4.7-4.8 (m, 1H), 6.85 (s, 1H), 7.1 (d, 2H), 7.25 (d, 1H), 7.3 (s, 1H), 7.45-7.55 (m, 4H), 8.05 (s, 1H), 12.6 (s, 1H)
1d		456 ($M+H$) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 2.95 (s, 3H), 3.4-3.6 (br, 4H), 4.7 (t, 1H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (d, 1H), 7.3 (s, 1H), 7.4-7.5 (m, 3H), 7.55 (d, 1H), 12.6 (s, 1H)
1e		442 ($M+H$) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 3.35 (m, 2H), 3.5 (m, 2H), 4.7 (t, 1H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (d, 2H), 7.3 (s, 1H), 7.35 (d, 1H), 7.9 (d, 2H), 8.4 (t, 1H), 12.6 (s, 1H)
1f		510 ($M+H$) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 3.2 (m, 4H), 3.4 (m, 4H), 4.75 (m, 1H), 6.25 (s, 1H), 6.85 (s, 1H), 7.1 (d, 2H), 7.25 (d, 2H), 7.5 (s, 1H), 7.55 (d, 1H), 7.85 (d, 2H), 8.45 (t, 1H), 12.6 (s, 1H)
1g		469 ($M+H$) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 2.6 (d, 3H), 3.85 (d, 2H), 4.7-4.8 (m, 1H), 6.85 (s, 1H), 7.1 (d, 1H), 7.25 (d, 2H), 7.3 (s, 1H), 7.35 (d, 2H), 7.7-7.8 (br, 1H), 7.95 (d, 2H), 8.65 (t, 1H), 12.7 (s, 1H)

1h		496 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.15-1.2 (m, 2H), 1.3 (d, 6H), 1.55-1.6 (d, 2H), 1.5-1.6 (m, 1H), 3.15 (t, 2H), 3.35 (t, 2H), 3.8 (d, 2H), 4.75 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (d, 2H), 7.45 (s, 1H), 7.5 (d, 1H), 7.9 (d, 2H), 8.45 (t, 1H), 12.6 (s, 1H)
1i		482 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 1.35 (br, 2H), 1.7-1.8 (br, 2H), 2.65 (s, 6H), 3.15 (t, 2H), 3.7 (m, 2H), 4.7 (t, 1H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (d, 1H), 7.3 (s, 1H), 7.4 (d, 1H), 7.5 (s, 1H), 7.55 (d, 1H), 12.6 (s, 1H)
1j		511 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 2.4 (m, 6H), 3.5 (m, 6H), 4.4 (t, 1H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (d, 1H), 7.3 (s, 1H), 7.4 (d, 2H), 7.5 (s, 1H), 7.55 (d, 1H), 12.6 (s, 1H)
1k		509 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 1.5-1.6 (br, 2H), 1.65-1.8 (br, 2H), 2.05 (s, 3H), 2.75 (br, 1H), 2.8 (s, 3H), 3.3 (br, 4H), 4.7-4.8 (m, 1H), 6.7 (s, 1H), 7.0 (d, 1H), 7.05 (d, 2H), 7.3 (s, 1H), 7.4 (m, 3H), 7.5 (s, 1H), 12.6 (s, 1H)
1l		506 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 2.0 (dt, 2H), 3.3 (t, 2H), 4.1 (t, 2H), 4.8 (m, 1H), 6.9 (s, 1H), 6.95 (s, 1H), 7.2 (d, 2H), 7.25 (s, 1H), 7.3 (d, 2H), 7.55 (s, 1H), 7.6 (d, 1H), 7.7 (s, 1H), 7.95 (d, 2H), 8.55 (t, 1H), 12.6 (s, 1H)
1m		535 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 1.65 (s, 4H), 1.8 (br, 2H), 2.2 (br, 1H), 3.0 (br, 1H), 3.3 (s, 9H), 4.7-4.8 (m, 1H), 6.7 (s, 1H), 6.9 (s, 1H), 7.05 (d, 2H), 7.3 (s, 1H), 7.35 (d, 1H), 7.4 (d, 2H), 7.5 (s, 1H), 12.6 (s, 1H)
1n[#]		426 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 2.95 (s, 6H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (d, 1H), 7.3 (d, 1H), 7.4-7.55 (m, 4H), 12.7 (s, 1H)

1o^{s#}		412 (M+H) ⁺	
1p^s		456 (M+H) ⁺	
1q^s		452 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 0.7-1.15 (m, 5H), 1.2 (d, 6H), 3.0 (t, 2H), 4.8 (m, 1H), 6.7 (s, 1H), 7.0 (d, 2H), 7.15 (d, 2H), 7.35 (s, 1H), 7.4 (d, 1H) 7.8 (d, 2H), 8.4 (t, 1H), 12.7 (s, 1H)
1r^s		502 (M+H) ⁺	
1s^s		507 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 1.8-1.9 (dt, 2H), 2.1-2.2 (t, 2H), 3.3-3.4 (m, 6H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (m, 2H), 7.5 (s, 1H), 7.5 (d, 1H) 7.85 (d, 2H), 8.45 (t, 1H), 12.6 (s, 1H)
1t^s		454 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 3.7-4.5 (br, 5H), 4.7-4.8 (m, 1H), 5.7 (d, 1H), 6.8 (s, 1H), 7.05 (d, 2H), 7.25 (d, 1H), 7.3 (d, 1H), 7.5 (s, 1H), 7.55 (d, 1H) 7.65 (d, 2H), 12.7 (s, 1H)
1u^s		466 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 3.5 (br, 4H), 3.6 (br, 4H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (d, 1H), 7.3 (d, 1H), 7.4-7.55 (m, 4H), 12.6 (s, 1H)

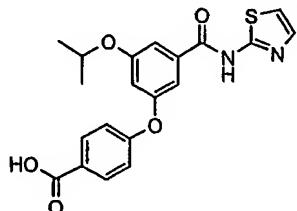
1v^s		509 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 2.0 (s, 3H), 3.4-3.6 (br, 8H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.1 (d, 2H), 7.25 (d, 1H), 7.3 (d, 1H), 7.4-7.55 (m, 4H), 12.7 (s, 1H)
1w^s		495 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 1.5-1.6 (m, 2H), 1.7-1.8 (m, 2H), 1.9-2.0 (m, 2H), 2.1 (s, 3H), 2.7-2.8 (d, 2H), 3.6-3.75 (br, 1H), 4.7-4.8 (m, 1H), 6.7 (s, 1H), 7.05 (d, 1H), 7.1 (d, 2H), 7.25 (s, 1H), 7.4 (d, 1H), 7.5 (s, 1H), 7.87 (d, 2H), 8.2 (d, 1H), 12.7 (s, 1H)
1x^s		478 (M+H) ⁺	
1y[*]		438 (M+H) ⁺	¹ H NMR δ (CDCl ₃): 1.35 (d, 6H), 2.36 (m, 2H), 4.20-4.38 (m, 4H), 4.58 (m, 1H), 6.78 (m, 1H), 7.00 (m, 3H), 7.17 (m, 1H), 7.27 (m, 2H), 7.63 (d, 2H)
1z		495 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 1.8 (d, 2H), 2.15 (d, 3H), 2.5 (d, 2H), 3.2-3.6 (m, 6H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.05 (d, 2H), 7.2 (d, 1H), 7.3 (s, 1H), 7.4 (d, 2H), 7.5 (s, 1H), 7.55 (d, 1H)

^sEDAC used as coupling reagent in place of HATU.

*Ethyl acetate was used as eluant.

= reference example

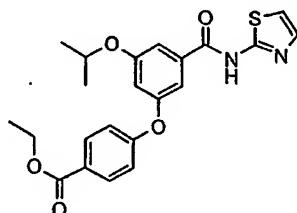
5 The required acid for Example 1 was prepared as described below:

4-({3-[(1-Methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenyl}oxy)benzoic acid

A solution of ethyl 4-({3-[(1-methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenyl}oxy)benzoate (2.5 g) in THF (100 mL) was added to a solution of lithium hydroxide

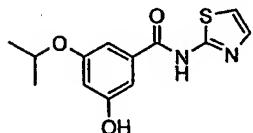
5 monohydrate (1.3 g) in water (50 mL). The mixture was stirred at ambient temperature for 16 hours and the THF removed *in vacuo*. The aqueous layer was acidified with 1M hydrochloric acid (30 mL), the solid precipitate filtered off, washed with water and dried *in vacuo* to give the desired compound (2.22 g).

¹H NMR δ (d₆-DMSO): 1.3 (d, 6H), 4.7-4.8 (m, 1H), 6.9 (t, 1H), 7.1 (d, 2H), 7.25 (d, 1H),
10 7.35 (s, 1H), 7.5 (s, 1H), 7.55 (d, 1H), 7.95 (d, 2H), 12.75 (s, 1H); *m/z* 399 (M+H)⁺

Ethyl 4-({3-[(1-methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenyl}oxy) benzoate

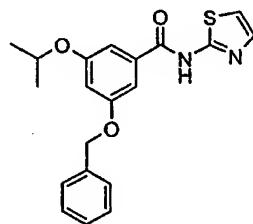
A solution of 3-hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide (3.06g), 4-
15 ethoxycarbonylphenylboronic acid (3.0g), copper (II) acetate (3.0g), triethylamine (7.6 mL) and freshly activated 4Å molecular sieves (12g) in DCM (170 mL) was stirred at ambient temperature and under ambient atmosphere for 2 days. The reaction mixture was filtered through diatomaceous earth, washed with DCM (2 x 50 mL), the DCM removed *in vacuo* and the residual oil partitioned between ethyl acetate (150 mL) and 1M hydrochloric acid (100 mL). The ethyl acetate layer was separated, washed with aqueous sodium hydrogen carbonate solution and brine, dried (MgSO₄), and evaporated to a residue which was chromatographed on silica with 20% ethyl acetate in isohexane as eluant to give the desired compound (2.64 g).

¹H NMR δ (CDCl₃): 1.3 (d, 6H), 1.35 (t, 3H), 4.35 (q, 2H), 4.5-4.6 (m, 1H), 6.8 (s, 1H), 6.95 (s, 1H), 7.0 (d, 2H), 7.15 (s, 1H), 7.2 (s, 1H), 7.3 (d, 1H), 8.05 (d, 2H); *m/z* 427 (M+H)⁺

3-Hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide

3-[(1-Methylethyl)oxy]-5-[(phenylmethyl)oxy]-N-1,3-thiazol-2-ylbenzamide (11.2 g) was dissolved in trifluoroacetic acid (60 mL) and treated with thioanisole (17.8 mL). The mixture 5 was left to stir at ambient temperature for 18 hours before the trifluoroacetic acid was removed *in vacuo*. The residues were treated with isohexane (100 mL) and the solid filtered off, before being washed with further isohexane (2 x 20 mL). The solid was dissolved in ethyl acetate (200 mL) and washed with aqueous saturated sodium hydrogen carbonate solution (100 mL). The organics were washed with water (100 mL) and brine (100 mL), and dried 10 (MgSO_4) before evaporation *in vacuo* to afford a solid which was washed with isohexane (200 mL) and dried *in vacuo* to give the desired compound (7.18 g).

$^1\text{H NMR}$ δ (d_6 -DMSO): 1.27 (d, 6H), 4.55 (m, 1H), 6.49 (m, 1H), 7.02 (s, 1H), 7.14 (s, 1H), 7.25 (d, 1H), 7.54 (d, 1H), 9.73 (s, 1H), 12.44 (s, 1H); m/z 279 ($\text{M}+\text{H})^+$, 277 ($\text{M}-\text{H}$)

15 3-[(1-Methylethyl)oxy]-5-[(phenylmethyl)oxy]-N-1,3-thiazol-2-ylbenzamide

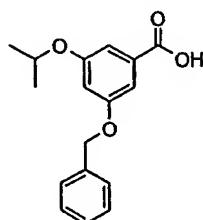
To a solution of 3-[(1-methylethyl)oxy]-5-[(phenylmethyl)oxy]benzoic acid (20 g) in DCM (400 mL), cooled to 0°C was slowly added oxalyl chloride (12.2 mL) and DMF (0.4 mL), with stirring. The mixture was allowed to warm to ambient temperature and stirred for a further 16 20 hours, following which the organics were removed *in vacuo*, and the residues azeotroped with toluene (100 mL). The crude material was dissolved in DCM (200 mL) and slowly added to a stirred suspension of 2-aminothiazole (10.5 g) and diisopropylethylamine (24.3 mL), in DCM (200 mL). The mixture was stirred at ambient temperature for 70 hours, before the organics were removed *in vacuo*. The residues were dissolved in ethyl acetate (300 mL) and washed 25 with 1M aqueous hydrochloric acid (300 mL). The aqueous layer was extracted with further ethyl acetate (300 mL), and the combined organics washed with brine (75 mL), and dried

(MgSO₄), before evaporation *in vacuo* to give the desired compound (28 g) which was used without further purification.

¹H NMR δ (d₆-DMSO): 1.27 (d, 6H), 4.70 (m, 1H), 5.15 (s, 2H), 6.77 (m, 1H), 7.27 (m, 2H), 7.33-7.47 (brm, 6H), 7.55 (d, 1H); m/z 369 (M+H)⁺, 367 (M-H)⁻;

5 The ¹H NMR spectrum also contained signals consistent with a small amount of ethyl acetate.

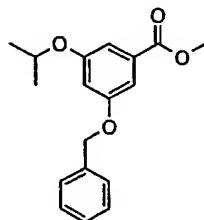
3-[(1-Methylethyl)oxy]-5-[(phenylmethyl)oxy]benzoic acid



To a solution of methyl 3-[(1-methylethyl)oxy]-5-[(phenylmethyl)oxy]benzoate (37 g) in a 1:1 mixture of THF:methanol (300 mL) was added 4M sodium hydroxide solution (150 mL). The mixture was refluxed for 45 minutes, following which the organics were removed *in vacuo*. The aqueous was acidified to pH4 with hydrochloric acid (2M), and extracted with ethyl acetate. The organics were combined, washed with water and brine, dried (MgSO₄) and concentrated *in vacuo* to give the desired compound (33.5 g), which was used without further purification.

¹H NMR δ (d₆-DMSO): 1.26 (d, 6H), 4.59-4.69 (m, 1H), 5.15 (s, 2H), 6.80 (app t, 1H), 7.04 (m, 1H), 7.12 (m, 1H), 7.33 (app t, 1H), 7.40 (t, 2H), 7.46 (d, 2H), 12.95 (s, 1H)

Methyl 3-[(1-methylethyl)oxy]-5-[(phenylmethyl)oxy]benzoate



20

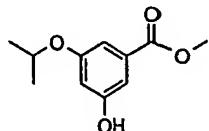
To a solution of methyl 3-hydroxy-5-[(1-methylethyl)oxy]benzoate (25 g) in DMF (250 mL) was added anhydrous potassium carbonate (297 mmol), and benzyl bromide (143 mmol). The mixture was stirred at 60°C for 5 hours, then cooled to room temperature. The solvent was removed *in vacuo* and the residue partitioned between ethyl acetate and water. The organics

were combined and washed with further water, brine, dried (MgSO_4) and concentrated *in vacuo* to give the desired compound (37 g) which was used without further purification.

^1H NMR δ (d₆-DMSO): 1.26 (d, 6H), 3.84 (s, 3H), 4.61-4.70 (m, 1H), 5.12 (s, 2H), 6.84 (t, 1H), 7.05 (app t, 1H), 7.12-7.15 (m, 1H), 7.31-7.37 (m, 1H), 7.40 (t, 2H), 7.46 (d, 2H)

5

Methyl 3-hydroxy-5-[(1-methylethyl)oxy]benzoate

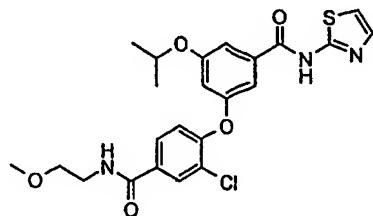


To a stirred solution of methyl 3,5-dihydroxybenzoate (0.1 mol) in DMF (180 mL) was added powdered potassium carbonate (0.2 mol) and 2-iodopropane (0.1 mol), and the resulting mixture stirred at ambient temperature for 16 hours. The reaction mixture was poured into water (1000 mL) and the mixture extracted with ether. The extracts were combined and washed sequentially with water (twice) and brine; the solution was dried (MgSO_4), filtered and evaporated *in vacuo* to give the crude product as a pale yellow oil (12.6 g). This was treated with toluene (40 mL) and allowed to stand overnight. The insoluble material (starting phenol) was removed by filtration, and the filtrate evaporated *in vacuo*. The resulting oil was chromatographed (2 x 90 g Biotage silica cartridges), eluting with hexane containing ethyl acetate (10% increasing to 15% v/v). The title compound was obtained as an oil (25% yield), which was identical by tlc to a sample prepared by a similar procedure.

^1H NMR δ (d₆-DMSO): 1.2 (d, 6H), 3.8 (s, 3H), 4.5 – 4.6 (hept, 1H), 6.55 (m, 1H), 7.85 (m, 1H), 7.95 (m, 1H), 9.8 (s, 1H)

20

Example 2: 3-Chloro-4-{3-[(1-methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenoxy}-N-(2-methoxyethyl)benzamide



25

To a suspension of 3-chloro-4-{3-[(1-methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenoxy}benzoic acid (107 mg), HATU (122 mg) and 2-methoxyethylamine (38 mg) in

DMF (2 mL) was added diisopropylethylamine (0.11 mL) and the mixture stirred at ambient temperature for 1 hour. Water (30 mL) was added and the mixture extracted with ethyl acetate (3 x 15 mL). The combined organic extracts were washed with brine, dried (MgSO_4), and evaporated to a residue which was chromatographed on silica with ethyl acetate as eluant
 5 to give the desired compound (85 mg).

^1H NMR δ (d_6 -DMSO): 1.3 (d, 6H), 3.25 (s, 3H), 3.4 (m, 4H), 4.7-4.8 (m, 1H), 6.85 (d, 1H), 7.2 (m, 1H), 7.5 (s, 1H), 7.55 (d, 1H), 7.8 (m, 1H), 8.05 (dd, 1H), 8.6 (t, 1H); m/z 486 ($\text{M}+\text{H}$)⁺.

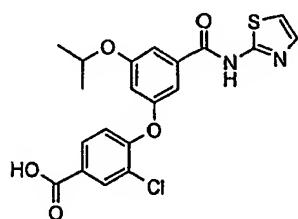
10 In a similar manner, Example 2a was also prepared:

Example	Structure	m/z	NMR
2a		460 ($\text{M}+\text{H}$) ⁺	^1H NMR δ (d_6 -DMSO): 1.3 (d, 6H), 2.95 (s, 6H), 4.7-4.8 (m, 1H), 6.8 (s, 1H), 7.2 (m, 2H), 7.25 (d, 1H), 7.4 (dd, 1H), 7.5 (s, 1H), 7.55 (d, 1H), 7.65 (s, 1H), 12.6 (s, 1H)

The required acid for Example 2 was prepared as described below:

3-Chloro-4-(3-[(1-methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenyl)oxybenzoic

15 acid



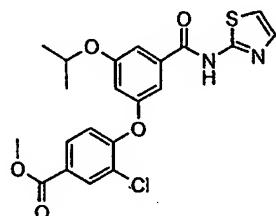
A solution of methyl 3-chloro-4-(3-[(1-methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenyl)oxybenzoate (950 mg) in THF (30 mL) was added to a solution of lithium hydroxide monohydrate (237 mg) in water (15 mL). The mixture was stirred at ambient

20 temperature for 16 hours and the THF removed *in vacuo*. The aqueous layer was acidified with 1M hydrochloric acid (5.3 mL), the solid precipitate filtered off, washed with water and dried *in vacuo* to give the desired acid (880 mg).

¹H NMR δ (d₆-DMSO): 1.3 (d, 6H), 4.7-4.8 (m, 1H), 6.9 (t, 1H), 7.15 (d, 1H), 7.25 (d, 2H), 7.5 (d, 1H), 7.55 (s, 1H), 7.9 (d, 1H), 8.05 (d, 1H), 12.75 (s, 1H)

Methyl 3-chloro-4-(3-[(1-methylethyl)oxy]-5-[(1,3-thiazol-2-ylamino)carbonyl]phenyl)oxy)benzoate

5 oxy)benzoate



To a solution of 3-hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide (208 mg) and methyl 3-chloro-4-fluorobenzoate (141 mg) in acetonitrile (5 mL) was added potassium carbonate (104 mg) and the stirred mixture heated at 160°C in a 'Smith Creator Microwave'

10 for 30 minutes. The mixture allowed to return to ambient temperature and pressure, the acetonitrile evaporated, and the residue partitioned between ethyl acetate (50 mL) and water (20 mL). The organic layer was separated, washed with brine, dried (MgSO₄), and evaporated to a residue which was chromatographed on silica, eluting with 30% ethyl acetate in isohexane, to give the desired ester (178 mg).

15 ¹H NMR δ (CDCl₃): 1.3 (d, 6H), 3.9 (s, 3H), 4.5-4.6 (m, 1H), 6.75 (t, 1H), 6.95 (d, 1H), 7.0 (d, 1H), 7.1 (s, 1H), 7.2 (m, 1H), 7.3 (s, 1H), 7.9 (dd, 1H), 8.05 (d, 1H); m/z 447 (M+H)⁺

The synthesis of 3-hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide is described above in Example 1.

20

Example 3: General Procedure for Preparation of Halogenated Sulphonamides

To a solution of the appropriate amine (1.8 mmol) in DCM (2 mL), was added the sulphonyl chloride (0.72 mmol) in DCM (2 mL), and the resulting mixture stirred for 18 hours. The mixture was treated with 1M aqueous hydrochloric acid (4 mL) and the organics separated. Evaporation *in vacuo* gave the crude fluorosulphonamide which was used without further purification.

To a solution of the crude fluorosulphonamide (7.2 mmol) in acetonitrile (3 mL), was added 3-hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide (0.36 mmol) and potassium carbonate (1.8 mmol). The mixture was heated to 170°C in a 'Smith Creator Microwave' for 100 minutes, before being filtered and the resultant organics evaporated *in vacuo*. The residues were then chromatographed on a Redisep (12 g, SiO₂) cartridge using an Isco Optix chromatography system, eluting with 30 to 100% ethyl acetate in isohexane, and evaporated *in vacuo* to afford the desired compound.

Using a similar procedure to that described above, Examples 3a-3e were prepared from 3-hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide:

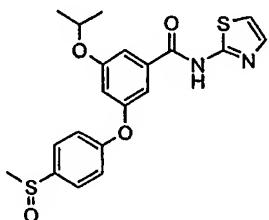
Example	Structure	<i>m/z</i>	NMR
3a		452 (M+H) ⁺ 450 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.28 (d, 6H), 4.70-4.80 (m, 1H), 6.93 (m, 1H), 7.26 (m, 2H), 7.37 (t, 1H), 7.47 (s, 2H), 7.54 (m, 2H), 7.68 (d, 1H), 7.80 (dd, 1H), 12.64 (s, 1H)
3b		496, 498 (M+H) ⁺ 494, 496 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.30 (d, 6H), 2.64 (s, 6H), 4.72-4.82 (m, 1H), 6.97 (m, 1H), 7.20-7.28 (m, 2H), 7.36 (m, 1H), 7.53 (m, 2H), 7.70 (dd, 1H), 7.92 (d, 1H), 12.64 (s, 1H)
3c		510, 512 (M+H) ⁺ 508, 510 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 0.98 (d, 6H), 1.30 (d, 6H), 4.68-4.79 (m, 2H), 6.92 (s, 1H), 7.21-7.31 (m, 3H), 7.53 (m, 2H), 7.66 (d, 2H), 7.76 (dd, 1H), 7.97 (m, 1H), amide NH not seen
3d ^s		551, 553 (M+H) ⁺ 549, 551 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.30 (d, 6H), 2.13 (s, 3H), 2.34 (s, 4H), 2.93 (s, 4H), 4.72-4.81 (m, 1H), 6.96 (s, 1H), 7.20-7.30 (m, 2H), 7.36 (s, 1H), 7.54 (s, 1H), 7.65-7.78 (m, 2H), 7.90 (m, 1H), amide NH not seen

3e		486, 488 ($M+H$) ⁺ 484, 486 ($M-H$) ⁻	^1H NMR δ (d_6 -DMSO): 1.29 (d, 6H), 4.70-4.82 (m, 1H), 6.97 (s, 1H), 7.26 (m, 2H), 7.30 (s, 1H), 7.47 (d, 1H), 7.54 (m, 2H), 7.73 (s, 1H), 7.92 (d, 1H), amide NH not seen
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⁵The requisite sulphonamide for this example was prepared using a 1:1 ratio of amine:sulphonyl chloride, and isolated by treatment with 1M aqueous sodium hydroxide. The synthesis of 3-hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide is described in Example 1 above.

5

Example 4: 3-(1-Methylethyl)oxy-5-[4-(1,3,4-oxadiazol-2-yl)phenoxy]-N-1,3-thiazol-2-ylbenzamide



- 10 A solution of 3-hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide (280 mg), 4-(methanesulphinyl)benzeneboronic acid (368 mg), copper (II) acetate (363 mg), triethylamine (0.7 mL) and freshly activated 4 Å molecular sieves (1.4 g) in DCM (10 mL), was stirred at ambient temperature and under ambient atmosphere for 3 days. The reaction mixture was filtered through diatomaceous earth, washed with DCM (2 x 10 mL), the DCM removed *in vacuo* and the residual oil partitioned between ethyl acetate (50 mL) and 1M hydrochloric acid (35 mL). The ethyl acetate layer was separated, washed sequentially with saturated aqueous sodium hydrogen carbonate and brine, dried (MgSO_4) and evaporated to a residue which was chromatographed on silica with 0-100% ethyl acetate in isohexane as eluant gave the desired compound (180 mg).
- 15 20 ^1H NMR δ (d_6 -DMSO): 1.28 (d, 6H), 2.74 (s, 3H), 4.74 (m, 1H), 6.86 (m, 1H), 7.20-7.33 (m, 4H), 7.50 (m, 1H), 7.53 (d, 1H), 7.72 (d, 2H), 12.62 (bs, 1H); m/z 417 ($M+H$)⁺

The following examples were synthesised in an analogous fashion:-

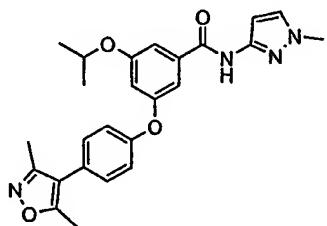
Example	Structure	m/z	NMR
4a		415 (M+H) ⁺ 413 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.20 (t, 3H), 1.29 (d, 6H), 2.94 (m, 2H), 4.73 (m, 1H), 6.77 (m, 1H), 7.05 (d, 2H), 7.20 (m, 1H), 7.26 (m, 1H), 7.37 (d, 2H), 7.45 (s, 1H), 7.53 (d, 1H), amide NH not seen
4b ^s		423 (M+H) ⁺ 421 (M-H) ⁻	¹ H NMR δ (CDCl ₃): 1.35 (d, 6H), 4.55 (m, 1H), 6.80 (m, 1H), 6.95 (m, 1H), 7.12 (d, 2H), 7.18 (m, 1H), 7.22 (m, 1H), 7.30 (m, 1H), 8.08 (d, 2H), 8.45 (s, 1H)

^sRequired further chromatography, eluting with 0-2% methanol in DCM.

The synthesis of 3-hydroxy-5-[(1-methylethyl)oxy]-N-1,3-thiazol-2-ylbenzamide is described in Example 1 above.

5

Example 5: 3-[4-(3,5-Dimethylisoxazol-4-yl)phenoxy]-5-(1-methylethyl)oxy-N-(1-methyl-1H-pyrazol-3-yl)benzamide

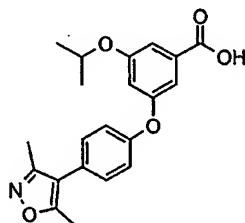


- 10 To a stirred solution of 3-[(4-(3,5-dimethylisoxazol-4-yl)phenyl)oxy]-5-[(1-methylethyl)oxy]benzoic acid (0.3 mmol) in DCM (2 mL) was added oxalyl chloride (50 µl) and a drop of DMF. The reaction was stirred overnight at room temperature, then evaporated in *vacuo*. The resulting acid chloride was dissolved in DCM (1 mL), and added to a solution of 1-methyl-1*H*-pyrazol-3-amine (0.38 mmol) and diisopropylethylamine (0.9 mmol) in DCM (2 mL).
- 15 The reaction was stirred at room temperature for 48 hours. The reaction mixture was diluted with DCM, and washed twice with 2M hydrochloric acid, then with saturated aqueous sodium hydrogen carbonate and brine. The solution was dried (MgSO₄), filtered, and evaporated to yield the product (84% yield).

¹H NMR δ (CDCl₃): 1.35 (d, 6H), 2.30 (s, 3H), 2.40 (s, 3H), 3.80 (s, 3H), 4.60 (m, 1H), 6.75 (m, 1H), 6.80 (m, 1H), 7.10 (m, 3H), 7.25 (br m, 4H), 8.70 (br s, 1H); *m/z* 447 (M+H)⁺

The preparation of 3-{{[4-(3,5-dimethylisoxazol-4-yl)phenyl]oxy}-5-[(1-methylethyl)oxy]benzoic acid is described below:

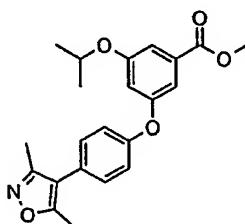
3-{{[4-(3,5-Dimethylisoxazol-4-yl)phenyl]oxy}-5-[(1-methylethyl)oxy]benzoic acid}



To a stirred solution of methyl 3-{{[4-(3,5-dimethylisoxazol-4-yl)phenyl]oxy}-5-[(1-methylethyl)oxy]benzoate (0.31 mmol) in THF (2 mL) was added lithium hydroxide (0.62 mmol) and water (0.35 mL). The reaction was stirred overnight at room temperature, before the addition of further lithium hydroxide (0.31 mmol) and water (0.2 mL). The reaction was stirred at room temperature for a further 3 hours, acidified with 2M hydrochloric acid and partitioned between water and ethyl acetate. The layers were separated and the aqueous layer reextracted with ethyl acetate. The combined organic layers were washed with brine, dried (MgSO₄), filtered, and evaporated to yield the product (*ca.* 100% yield).

¹H NMR δ (CDCl₃): 1.35 (d, 6H), 2.30 (s, 3H), 2.45 (s, 3H), 4.60 (s, 1H), 6.85 (m, 1H), 7.10 (d, 2H), 7.22 (d, 2H), 7.35 (m, 1H), 7.40 (m, 1H); *m/z* 368 (M+H)⁺, 366 (M-H)⁻

20 Methyl 3-{{[4-(3,5-dimethylisoxazol-4-yl)phenyl]oxy}-5-[(1-methylethyl)oxy]benzoate}

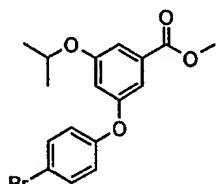


Methyl 3-[(4-bromophenyl)oxy]-5-[(1-methylethyl)oxy]benzoate (0.74 mmol) and 3,5-dimethylisoxazole-4-boronic acid (0.81 mmol) were suspended in a 1:1 mixture of dimethoxyethane and 2M sodium carbonate (6 mL). The mixture was degassed, before the addition of tetrakis(triphenylphosphine)palladium (0.015 mmol). The mixture was again

degassed, and stirred at 80°C, then at room temperature overnight. The reaction was filtered through diatomaceous earth then evaporated in *vacuo*. The residual oil was partitioned between ethyl acetate and 2M sodium hydroxide. The ethyl acetate layer was separated, washed with brine, dried ($MgSO_4$), and evaporated to a residue which was chromatographed 5 on silica with 10% ethyl acetate in isohexane as eluant to give the desired ester (43% yield).

1H NMR δ ($CDCl_3$): 1.35 (d, 6H), 2.25 (s, 3H), 2.45 (s, 3H), 3.90 (s, 3H), 4.60 (m, 1H), 6.80 (m, 1H), 7.10 (d, 2H), 7.25 (br m, 3H), 7.35 (br s, 1H); *m/z* 382 ($M+H$)⁺

Methyl 3-[*(4*-bromophenyl)oxy]-5-[*(1*-methylethyl)oxy]benzoate



10

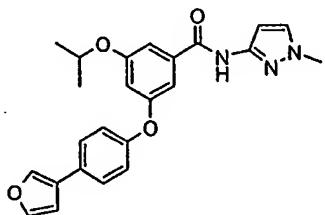
A solution of methyl 3-hydroxy-5-[*(1*-methylethyl)oxy]benzoate (0.024 mol), 4-bromophenyl boronic acid (0.048 mol), copper (II) acetate (0.048 mol), triethylamine (0.12 mol) and freshly activated 4Å molecular sieves (25 g) in DCM (500 mL) was stirred at ambient temperature and under ambient atmosphere for 7 days. The reaction mixture was filtered, the DCM 15 removed in *vacuo*, and the residual oil partitioned between ethyl acetate and 2M hydrochloric acid. The ethyl acetate layer was separated, washed sequentially with saturated aqueous sodium hydrogen carbonate, brine, dried ($MgSO_4$), and evaporated to a residue which was chromatographed on silica, eluting with 10-40% ethyl acetate in isohexane, to give the desired ester (56% yield).

20 1H NMR δ (d_6 -DMSO): 1.25 (d, 6H), 3.80 (s, 3H), 4.65 (m, 1H), 6.87 (m, 1H), 6.97 (m, 1H), 7.03 (d, 2H), 7.20 (m, 1H), 7.55 (d, 2H); *m/z* 367 ($M+H$)⁺

The synthesis of methyl 3-hydroxy-5-[*(1*-methylethyl)oxy]benzoate is described above in Example 1.

25

Example 6: 3-[(4-Furan-3-ylphenyl)oxy]-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide



In a similar fashion to that described above for **Example 5**, 3-[(4-furan-3-ylphenyl)oxy]-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide was prepared from 3-[(4-furan-3-ylphenyl)oxy]-5-[(1-methylethyl)oxy]benzoic acid.

¹H NMR δ (CDCl₃): 1.35 (d, 6H), 3.75 (s, 3H), 4.55 (m, 1H), 6.65 (s, 1H), 6.70 (m, 1H), 6.80 (m, 1H), 6.98 (s, 1H), 7.02 (d, 2H), 7.12 (s, 1H), 7.25 (m, 1H), 7.45 (m, 3H), 7.70 (s, 1H), 8.60 (br s, 1H); m/z 418 (M+H)⁺

10

3-[(4-Furan-3-ylphenyl)oxy]-5-[(1-methylethyl)oxy]benzoic acid was prepared from methyl 3-[(4-bromophenyl)oxy]-5-[(1-methylethyl)oxy]benzoate in an analogous manner to 3-[(4-(3,5-dimethylisoxazol-4-yl)phenyl)oxy]-5-[(1-methylethyl)oxy]benzoic acid described in **Example 5**.

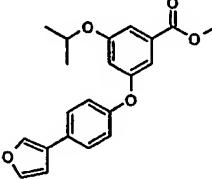
15

3-[(4-Furan-3-ylphenyl)oxy]-5-[(1-methylethyl)oxy]benzoic acid

Structure	NMR	m/z
	¹ H NMR δ (CDCl ₃): 1.35 (d, 6H), 4.60 (m, 1H), 6.65 (m, 1H), 6.80 (m, 1H), 7.05 (d, 2H), 7.30 (m, 1H), 7.35 (m, 1H), 7.45 (m, 3H), 7.70 (s, 1H)	337 (M-H) ⁻

Methyl 3-[(4-furan-3-ylphenyl)oxy]-5-[(1-methylethyl)oxy]benzoate

20

Structure	NMR	m/z
	¹ H NMR δ (CDCl ₃): 1.35 (d, 6H), 3.90 (s, 3H), 4.55 (m, 1H), 6.65 (s, 1H), 6.72 (m, 1H), 7.02 (d, 2H), 7.22 (m, 1H), 7.30 (m, 1H), 7.45 (m, 3H), 7.70 (s, 1H)	353 (M+H) ⁺

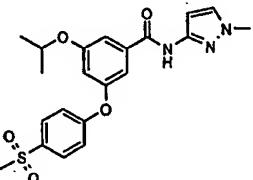
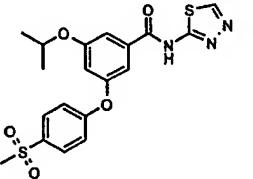
Example 7: General Procedure for Amide Synthesis – HATU Coupling

5 Diisopropylethylamine (2.5 equivalents) was added to a suspension of 3-{(1-methylethyl)oxy}-5-{[4-(methylsulfonyl)phenyl]oxy}benzoic acid (1 equivalent), HATU (1.25 equivalents) and amine (1.25 equivalents) in DMF (20 mL). The initial suspension dissolved into a dark orange solution. The resulting mixture was stirred at ambient temperature for 2 hours. The DMF was removed *in vacuo*, and the residue azeotroped with

10 toluene. Water was added and the mixture extracted with ethyl acetate. The extracts were combined and washed sequentially with 1M hydrochloric acid, saturated sodium hydrogen carbonate solution and brine. The solution was dried (MgSO₄), filtered, and evaporated *in vacuo* to give the crude product which was chromatographed, eluting with 50% ethyl acetate in isohexane, to give desired compound (40-70% yield).

15

Using the above method, Examples 7a-7c were prepared:

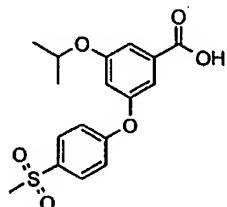
Example	Structure	m/z	NMR
7a		430 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.25 (d, 6H), 3.2 (s, 3H), 3.8 (s, 3H), 4.75 (m, 1H), 6.55 (d, 1H), 6.85 (s, 1H), 7.2 (d, 2H), 7.3 (s, 1H), 7.45 (s, 1H), 7.6 (d, 1H), 7.9 (d, 2H), 10.85 (br s, 1H)
7b		434 (M+H) ⁺ 432 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.25 (d, 6H), 3.2 (s, 3H), 4.75 (m, 1H), 7.0 (s, 1H), 7.2 (d, 2H), 7.4 (s, 1H), 7.6 (s, 1H), 7.95 (d, 1H), 9.2 (s, 1H), 13.10 (br s, 1H)

7c[#]		433 (M+H) ⁺ , 431 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.3 (d, 6H), 3.2 (s, 3H), 4.75 (m, 1H), 7.0 (s, 1H), 7.2 (d, 2H), 7.25 (s, 1H), 7.4 (s, 1H), 7.6 (s, 1H), 7.65 (s, 1H), 7.95 (d, 1H)
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[#]Example 7c may be crystallised by allowing isohexane to vapour diffuse into a solution of the compound in ethylacetate, in a closed system, with subsequent slow evaporation of the mixture at room temperature over 4 days, mpt 145-147°C

5 The required acid for the synthesis of Examples 7a-7c was prepared as described below:

3-((1-Methylethyl)oxy)-5-((4-(methylsulfonyl)phenyl)oxy)benzoic acid

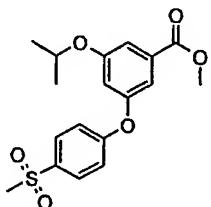


A solution of methyl 3-((1-methylethyl)oxy)-5-((4-(methylsulfonyl)phenyl)oxy)benzoate

10 (15.1 mmol) in THF (90 mL) was treated with a solution of 1M sodium hydroxide (37 mmol), and the reaction mixture stirred for 13 hours at ambient temperature. Most of the organic solvent was removed *in vacuo*, and the remaining solution was diluted with water (50 mL). The resulting aqueous solution was acidified to pH4 with 1M citric acid solution, and extracted with ethyl acetate (2 x 40 mL). The extracts were combined, washed with brine, 15 dried (MgSO₄), and evaporated to give the desired compound (82% yield).

¹H NMR δ (d₆-DMSO): 1.25 (d, 3H), 3.2 (s, 3H), 4.64 (m, 1H), 6.95 (s, 1H), 7.06 (s, 1H), 7.2(d, 2H), 7.25 (s, 1H), 7.95 (d, 2H); *m/z* 349 (M-H)⁻

Methyl 3-((1-methylethyl)oxy)-5-((4-(methylsulfonyl)phenyl)oxy)benzoate



A suspension of methyl 3-hydroxy-5-[(1-methylethyl)oxy]benzoate (24 mmol), boronic acid (1.1 equivalents), copper (II) acetate (1.1 equivalents), triethylamine (5 equivalents) and freshly activated 4Å molecular sieves (31 g) in DCM (250 mL) was stirred at ambient temperature and under ambient atmosphere for 2 days. The reaction mixture was filtered, the

- 5 DCM removed *in vacuo* and the residual oil partitioned between ethyl acetate and 1-2M hydrochloric acid. The ethyl acetate layer was separated, washed with aqueous sodium hydrogen carbonate and brine, dried ($MgSO_4$), and evaporated to a residue which was chromatographed on silica (with 10-40% ethyl acetate in isohexane as eluant) to give the desired ester (64% yield).
- 10 1H NMR δ (d₆-DMSO): 1.25 (d, 3H), 3.2 (s, 3H), 4.64 (m, 1H), 6.95 (s, 1H), 7.06 (s, 1H), 7.2(d, 2H), 7.25 (s, 1H), 7.95 (d, 2H); *m/z* 365 (M+H)⁺

The synthesis of methyl 3-hydroxy-5-[(1-methylethyl)oxy]benzoate is described above in **Example 1.**

15

Example 8: General Procedure for Amide Synthesis – Phosphorus Oxychloride Coupling

- 20 Phosphorus oxychloride (0.75 mmol; 1.5 equivalents) was added dropwise to a stirred solution of 3-{(1-methylethyl)oxy}-5-{{[4-(methylsulfonyl)phenyl]oxy}benzoic acid (0.5 mmol) and the appropriate amine (1.25 equivalents) in pyridine (5 mL). The resulting mixture was stirred at ambient temperature for 4 hours. The pyridine was removed *in vacuo*, and the residue taken up in ethyl acetate. The mixture was washed sequentially with water, 1M citric acid and brine, dried ($MgSO_4$), filtered, and evaporated *in vacuo* to give the crude product,
- 25 which was chromatographed, eluting with 30-90% ethyl acetate in isohexane, to give the desired product (~20% yield).

Using the above method, Examples 8a & 8b were prepared:

Example	Structure	m/z	NMR
8a		427 (M+H) ⁺ 425 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.25 (d, 6H), 3.2 (s, 3H), 4.75 (m, 1H), 6.9 (s, 1H), 7.15 (m, 1H), 7.2 (d, 2H), 7.3 (s, 1H), 7.5 (s, 1H), 7.8 (app t, 1H), 7.95 (d, 2H), 8.15 (d, 1H), 8.4 (d, 1H), 10.8 (br s, 1H)
8b		428 (M+H) ⁺ 426 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.25 (d, 6H), 3.2 (s, 3H), 4.75 (m, 1H), 6.95 (s, 1H), 7.2 (d, 2H), 7.35 (s, 1H), 7.5 (s, 1H), 7.95 (d, 2H), 8.4 (d, 1H), 8.45 (d, 1H), 9.4 (d, 1H), 11.15 (br s, 1H)

The synthesis of 3-((1-methylethyl)oxy)-5-((4-(methylsulfonyl)phenyl)oxy)benzoic acid is described in Example 7 above.

5

Example 9: General Procedure for Amide Synthesis – Oxalyl Chloride Coupling

To a stirred solution of 3-((1-methylethyl)oxy)-5-((4-(methylsulfonyl)phenyl)oxy)benzoic acid (0.285 mmol) in dry DCM (2 mL), was added, dropwise under argon, oxalyl chloride (2 equivalents) and DMF (1 drop). The resulting solution was stirred at ambient temperature for 1-2 hours. The solvent was removed *in vacuo* and the crude mixture taken up in pyridine (2 mL) and added to the appropriate amine (2.2 equivalents). The reaction mixture was stirred at room temperature, or heated if necessary, and monitored by TLC and/or LCMS. The pyridine was removed *in vacuo*, and water and ethyl acetate added. The organic layer was washed sequentially with 1M citric acid and brine solution and dried ($MgSO_4$), concentrated *in vacuo*, and the residue chromatographed on silica (eluting with 30-90% ethyl acetate in isohexane) to give the desired product (typically 35-40% yield).

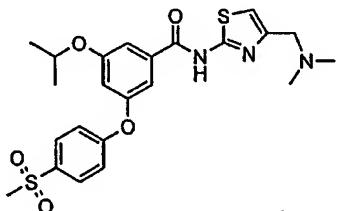
In a similar manner, Examples 9a-9c were prepared:-

Example	Structure	m/z	NMR
9a		431 (M+H) ⁺ 429 (M-H) ⁻	¹ H NMR δ (d ₆ -DMSO): 1.25 (d, 6H), 2.4 (s, 3H), 3.2 (s, 3H), 4.75 (m, 1H), 6.7 (s, 1H), 6.95 (s, 1H), 7.2 (d, 2H), 7.3 (s, 1H), 7.45 (s, 1H), 7.9 (d, 2H), 11.3 (br s, 1H)
9b		417 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.30 (dd, 6H), 3.23 (s, 3H), 4.78 (sept, 1H), 6.96 (s, 1H), 7.03 (s, 1H), 7.25 (d, 2H), 7.32 (s, 1H), 7.50 (s, 1H), 7.96 (d, 2H), 8.87 (s, 1H), 11.46 (s, 1H)
9c		500 (M+H) ⁺	¹ H NMR δ (d ₆ -DMSO): 1.33 (d, 6H), 3.23 (s, 3H), 4.78 (m, 1H), 6.77 (d, 1H), 7.01 (t, 1H), 7.26 (m, 3H), 7.42 (s, 1H), 7.60 (s, 1H), 7.98 (m, 3H), 13.22 (s, 1H)

The synthesis of 3-((1-methylethyl)oxy)-5-((4-(methylsulfonyl)phenyl)oxy)benzoic acid is described in Example 7 above.

5

Example 10: *N*-{4-[(Dimethylamino)methyl]-1,3-thiazol-2-yl}-3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]benzamide

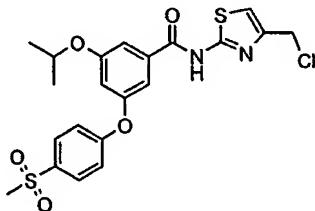


To a solution of *N*-{4-chloromethyl-1,3-thiazol-2-yl}-3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]benzamide (1.0 mmol) in THF (4 mL) was added dimethylamine in THF (10 mL of a 2M solution) and stirred at ambient temperature for 13 hours. The reaction mixture was concentrated *in vacuo* and the residue taken up in ethyl acetate. The mixture was washed sequentially with 1M sodium hydroxide and brine, dried (MgSO₄), and concentrated *in vacuo*. The residue was chromatographed, eluting with 20-80% ethyl acetate in isohexane, to give the desired compound (15% yield).

¹H NMR δ (d₆-DMSO): 1.3 (d, 6H), 2.2 (s, 6H), 3.2 (s, 3H), 3.4 (s, 2H), 4.75 (m, 1H), 6.9 (s, 1H), 7.0 (s, 1H), 7.2 (d, 2H), 7.35 (s, 1H), 7.55 (s, 1H), 7.95 (d, 1H); m/z 490 (M+H)⁺, 488 (M-H)⁻

5 The required chloromethylthiazole for Example 10 was prepared as described below:

N-{4-Chloromethyl-1,3-thiazol-2-yl}-3-(1-methylethyl)oxy-5-[4-(methylsulfonyl)phenoxy]benzamide



10 To a stirred solution of 3-{(1-methylethyl)oxy}-5-{[4-(methylsulfonyl)phenyl]oxy}benzoic acid (1.0 mmol) in DCM (10 mL) was added 1 drop of DMF and oxalyl chloride (2.0 mmol; 2.0 equivalents) dropwise, and the resulting mixture stirred at ambient temperature under argon for 2 hours. The reaction mixture was concentrated *in vacuo* and azeotroped with DCM. The residue was dissolved in DCM and 4-chloromethylthiazol-2-ylamine (1.0 mmol)

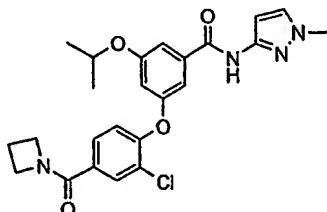
15 in DCM, diisopropylethylamine (2.5 mmol) and dimethylaminopyridine (0.1 mmol) added. The resulting mixture was stirred for 13 hours under argon at ambient temperature. The reaction was concentrated *in vacuo*, and chromatographed, eluting with 50-60% ethyl acetate in isohexane, to give the desired compound (53% yield).

¹H NMR δ (CDCl₃): 1.3 (d, 6H), 2.2 (s, 6H), 3.2 (s, 3H), 3.4 (s, 2H), 4.75 (m, 1H), 6.9 (s, 2H), 7.0 (s, 1H), 7.2 (d, 2H), 7.35 (s, 1H), 7.55 (s, 1H), 7.95 (d, 1H)

The synthesis of 3-{(1-Methylethyl)oxy}-5-{[4-(methylsulfonyl)phenyl]oxy}benzoic acid is described above in Example 7.

25 The preparation of 4-chloromethylthiazol-2-ylamine is described in the literature (*J. Indian Chem. Soc.*, 1960, 37, 241).

Example 11: 3-{[4-(Azetidin-1-ylcarbonyl)-2-chlorophenyl]oxy}-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide



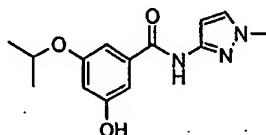
Potassium carbonate (300 mg, 2.18 mmol) was added to a mixture of 3-hydroxy-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide (300 mg, 0.81 mmol) and 1-[(3-chloro-4-fluorophenyl)carbonyl]azetidine (269 mg, 1.31 mmol) in DMF (5.0 mL) and the stirred mixture heated at 160°C in a ‘Smith Creator Microwave’ for 2 hours. The mixture was allowed to reach ambient temperature and pressure and reduced *in vacuo*. The residual oil was partitioned between ethyl acetate (50 mL) and water (50 mL). The ethyl acetate layer was separated, washed with brine, dried (MgSO_4), and evaporated to a residue which was chromatographed on silica, eluting with ethyl acetate, to give the desired compound (394 mg)

$^1\text{H NMR}$ δ (CDCl_3): 1.35 (d, 6H), 2.37 (quin, 2H), 3.80 (s, 3H), 4.20-4.20 (brm, 4H), 4.59 (m, 1H), 6.69 (m, 1H), 6.79 (m, 1H), 7.00 (m, 2H), 7.18 (m, 1H), 7.27 (m, 1H), 7.50 (d, 1H), 7.79 (s, 1H), 8.43 (brs, 1H); m/z 469, 471 ($\text{M}+\text{H})^+$

15

The required phenol for Example 11 was prepared as described below:-

3-Hydroxy-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide



20 3-[(1-Methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)-5-[(phenylmethyl)oxy]benzamide (51 g; 0.14 mol) was dissolved in methanol (500 mL) and THF (500 mL) and the flask evacuated and purged with argon (3 times). 10% Palladium on carbon (5.1 g) was added and the flask further evacuated and finally purged with hydrogen gas. The reaction mixture was stirred at ambient temperature for 20 hours. The reaction mixture was evacuated and purged with

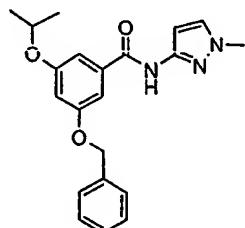
25 nitrogen (3 times). The catalyst was filtered off through celite, and the filtrate concentrated *in*

vacuo. Ethyl acetate was added and filtered to give the desired compound. (30.5 g) A second crop of material was obtained in the same way. (4.0 g)

^1H NMR δ (d_6 -DMSO): 1.30 (d, 6H), 3.78 (s, 3H), 4.68 (sept, 1H), 6.47 (m, 1H), 6.60 (s, 1H), 6.94 (s, 1H), 7.05 (s, 1H), 7.60 (s, 1H), 10.63 (s, 1H); m/z 276 ($M+\text{H}$)⁺

5

3-[(1-Methylethyl)oxy]-N-(1-methyl-1*H*-pyrazol-3-yl)-5-[(phenylmethyl)oxy]benzamide



DMF (2 drops) was added to a solution of 3-[(1-methylethyl)oxy]-5-[(phenylmethyl)oxy]benzoic acid (40.0 g, 0.14 mol) and oxalyl chloride (14.6 mL, 0.17 mol)

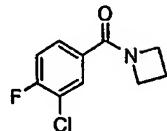
10 in DCM (700 mL) The mixture was stirred at ambient temperature for 4 hours and the DCM and excess oxalyl chloride evaporated *in vacuo*. The residual acid chloride was dissolved in DCM (300 mL) and added dropwise to 1-methyl-3-aminopyrazole (14.25 g, 0.147 mol) and triethylamine (41 mL, 0.29 mol) in DCM (300 mL), at 0°C. Stirred at ambient temperature for 24 hours. The DCM was evaporated *in vacuo*, and the residue partitioned between ethyl acetate (400 mL) and 1N hydrochloric acid (200 mL). The ethyl acetate layer was washed sequentially with saturated aqueous sodium hydrogen carbonate (200 mL) and brine (100 mL), dried (MgSO_4) and evaporated *in vacuo*. The residue was chromatographed on silica, eluting with a gradient of 50% ethyl acetate in isohexane, to give the desired compound (51 g)

15 ^1H NMR δ (CDCl_3): 1.30 (d, 6H), 3.61 (s, 3H), 4.50 (sept, 1H), 5.01 (s, 2H), 6.66 (m, 1H), 6.88 (m, 1H), 7.00 (m, 1H), 7.06 (m, 1H), 7.24 (m, 1H), 7.39 (m, 5H), 9.50 (s, 1H). m/z 366 ($M+\text{H}$)⁺

The preparation of 3-[(1-methylethyl)oxy]-5-[(phenylmethyl)oxy]benzoic acid was described in Example 1.

25

The preparation of 1-[(3-chloro-4-fluorophenyl)carbonyl]azetidine is described below:

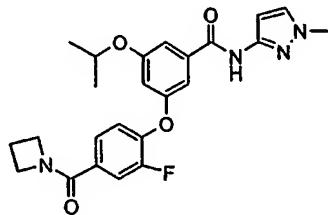
1-[(3-Chloro-4-fluorophenyl)carbonyl]azetidine

To a solution of 3-chloro-4-fluorobenzoic acid (1.74 g, 10.0 mmol) in DCM (50 mL) was added oxalyl chloride (1.05 mL, 12.0 mmol) and DMF (1 drop). The mixture was stirred at ambient temperature for 16 hours and the DCM and excess oxalyl chloride evaporated *in vacuo*. The residual acid chloride and azetidine hydrochloride (1.12 g, 12 mmol) were taken up in DCM (25 mL) and triethylamine (4.18 mL, 30 mmol) added to the mixture, which was stirred at ambient temperature for 2 hours. The DCM was evaporated *in vacuo*, and the residue partitioned between ethyl acetate (100 mL) and 1N hydrochloric acid (50 mL). The ethyl acetate layer was washed sequentially with saturated aqueous sodium hydrogen carbonate and brine, dried (MgSO_4), and evaporated. The residue was crystallized from ethyl acetate / isohexane to give the title compound (1.64 g).

^1H NMR δ (CDCl_3): 2.4 (m, 2H), 4.2-4.4 (m, 4H), 7.2 (m, 1H), 7.55 (m, 1H), 7.7 (m, 1H)

15

Example 12: 3-{[4-(Azetidin-1-yl)carbonyl]-2-fluorophenyl}oxy}-5-[(1-methylethyl)oxy]-
N-(1-methyl-1*H*-pyrazol-3-yl)benzamide

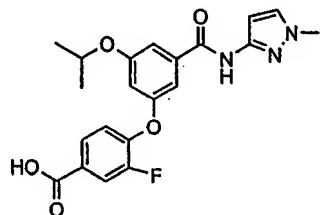


To a suspension of 3-fluoro-4-[(3-[(1-methylethyl)oxy]-5-[(1-methyl-1*H*-pyrazol-3-yl)amino]carbonyl}phenyl]oxy]benzoic acid (300 mg, 0.73 mmol), HATU (590 mg, 1.52 mmol) and azetidine hydrochloride (138 mg, 1.45 mmol) in DMF (5 mL), was added diisopropylethylamine (0.52 mL, 2.9 mmol) and the mixture stirred at ambient temperature for 24 hours. Water (30 mL) was added and the mixture extracted with ethyl acetate (3 x 15 mL). The combined organic extracts were washed with brine, dried (MgSO_4), and evaporated to a residue which was chromatographed on silica, eluting with ethyl acetate, to give the desired compound (190 mg).

¹H NMR δ (CDCl₃): 1.38 (d, 6H), 2.39 (m, 2H), 3.79 (s, 3H), 4.20-4.42 (m, 4H), 4.58 (sept, 1H), 6.70 (m, 1H), 6.79 (m, 1H), 7.00 (m, 1H), 7.07 (t, 1H), 7.16 (m, 1H), 7.27 (m, 1H), 7.41 (d, 1H), 7.51 (d, 1H), 8.44 (brs, 1H); m/z 453 (M+H)⁺

5 The required acid for Example 12 was prepared as described below:

3-Fluoro-4-[(3-[(1-methylethyl)oxy]-5-{[(1-methyl-1*H*-pyrazol-3-yl)amino]carbonyl}phenyl)oxy]benzoic acid



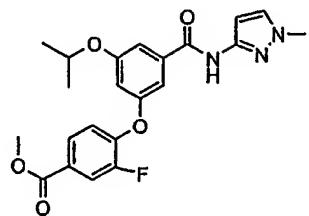
10 The mixture of methyl 3-fluoro-4-[(3-[(1-methylethyl)oxy]-5-{[(1-methyl-1*H*-pyrazol-3-yl)amino]carbonyl}phenyl)oxy]benzoate and 3-fluoro-4-[(3-[(1-methylethyl)oxy]-5-{[(1-methyl-1*H*-pyrazol-3-yl)amino]carbonyl}phenyl)oxy]benzoic acid isolated from the previous reaction (0.75 g, 1.76 mmol) was dissolved in THF (30 mL) and added to a solution of lithium hydroxide monohydrate (0.37 g, 8.8 mmol) in water (15mL). The mixture was stirred at

15 ambient temperature for 20 hours and the THF removed *in vacuo*. The aqueous layer was acidified with 1M hydrochloric acid (10 mL), the solid precipitate filtered off, washed with water and dried *in vacuo* to give the desired compound (0.57 g).

18 ¹H NMR δ (d₆-DMSO): 1.29 (d, 6H), 3.78 (s, 3H), 4.71 (sept, 1H), 6.55 (m, 1H), 6.83 (m, 1H), 7.22 (m, 2H), 7.41 (s, 1H), 7.57 (m, 1H), 7.81 (m, 2H), 10.83 (brs, 1H).

20 m/z 414 (M+H)⁺

Methyl 3-fluoro-4-[(3-[(1-methylethyl)oxy]-5-{[(1-methyl-1*H*-pyrazol-3-yl)amino]carbonyl}phenyl)oxy]benzoate



Potassium carbonate (500 mg, 3.64 mmol) was added to a mixture of 3-hydroxy-5-[(1-methylethyl)oxy]-N-(1-methyl-1*H*-pyrazol-3-yl)benzamide (500 mg, 1.82 mmol) and methyl-3,4-difluorobenzoate (370 mg, 2.18 mmol) in DMF (5.0 mL) and the stirred mixture heated at 160°C in a 'Smith Creator Microwave' for 2 hours. The mixture was allowed to reach

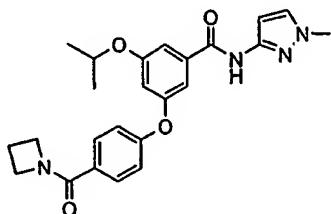
5 ambient temperature and pressure and reduced *in vacuo*. Water (10 mL) was added, taken to pH4 and extracted into ethyl acetate (2 x 15 mL), washed with brine (10 mL) and dried (MgSO_4) to give a mixture of the desired compound and 3-fluoro-4-[(3-[(1-methylethyl)oxy]-5-{{[(1-methyl-1*H*-pyrazol-3-yl)amino]carbonyl}phenyl}oxy]benzoic acid (0.75 g). This mixture was used without further purification.

10 m/z 428 and 414 ($\text{M}+\text{H}$)⁺

The preparation of 3-hydroxy-5-[(1-methylethyl)oxy]-N-(1-methyl-1*H*-pyrazol-3-yl)benzamide is described in Example 11.

15

Example 13: 3-{{[4-(Azetidin-1-ylcarbonyl)phenyl]oxy}-5-[(1-methylethyl)oxy]-N-(1-methyl-1*H*-pyrazol-3-yl)benzamide

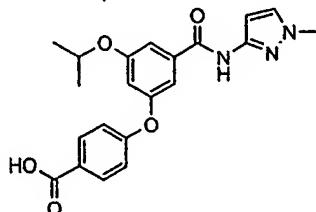


To a suspension of 4-[(3-[(1-methylethyl)oxy]-5-{{[(1-methyl-1*H*-pyrazol-3-yl)amino]carbonyl}phenyl}oxy]benzoic acid (300 mg, 0.63 mmol), HATU (600 mg, 1.60 mmol) and azetidine hydrochloride (150 mg, 1.52 mmol) in DMF (5 mL), was added diisopropylethylamine (0.56 mL, 3.04 mmol) and the mixture stirred at ambient temperature for 24 hours. Water (30 mL) was added and the mixture extracted with ethyl acetate (3 x 15 mL). The combined organic extracts were washed with brine, dried (MgSO_4), and evaporated to a residue which was chromatographed on silica, eluting with ethyl acetate, to give the desired compound (120 mg).

^1H NMR δ (d_6 -DMSO): 1.30 (d, 6H), 2.25 (m, 2H), 3.78 (s, 3H), 4.05 (m, 2H), 4.43 (m, 2H), 4.75 (sept, 1H), 6.58 (s, 1H), 6.81 (s, 1H), 7.09 (d, 2H), 7.23 (s, 1H), 7.44 (s, 1H), 7.60 (s, 1H), 7.69 (d, 2H), 10.80 (brs, 1H); m/z 435 ($\text{M}+\text{H}$)⁺

The required acid for Example 13 was prepared as described below:-

4-[(3-[(1-Methylethyl)oxy]-5-{{(1-methyl-1H-pyrazol-3-yl)amino]carbonyl}phenyl)oxy]benzoic acid



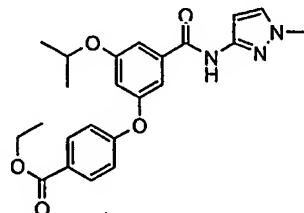
5

A solution of ethyl 4-[(3-[(1-methylethyl)oxy]-5-{{(1-methyl-1H-pyrazol-3-yl)amino]carbonyl}phenyl)oxy]benzoate (14.74 g, 0.035 mol) in THF (580 mL) was added to a solution of lithium hydroxide monohydrate (7.31 g, 0.175 mol) in water (290 mL). The mixture was stirred at ambient temperature for 48 hours and the THF removed *in vacuo*. The aqueous layer was acidified with 1M hydrochloric acid (10 mL), and the solid precipitate filtered off, washed with water and dried *in vacuo* to give the desired compound (12.6 g).

10 ^1H NMR δ (d_6 -DMSO): 1.28 (d, 6H), 3.78 (s, 3H), 4.71 (sept, 1H), 6.54 (m, 1H); 6.81 (m, 1H), 7.09 (d, 2H), 7.24 (s, 1H), 7.42 (s, 1H), 7.59 (m, 1H), 7.98 (d, 2H), 10.85 (brs, 1H), 12.80 (brs, 1H). m/z 396 (M+H)⁺

15

Ethyl 4-[(3-[(1-methylethyl)oxy]-5-{{(1-methyl-1H-pyrazol-3-yl)amino]carbonyl}phenyl)oxy]benzoate



Potassium carbonate (1g, 7.26 mmol) was added to a mixture of 3-hydroxy-5-[(1-methylethyl)oxy]-N-(1-methyl-1H-pyrazol-3-yl)benzamide (1 g, 3.63 mmol) and ethyl-4-fluorobenzoate (672 mg, 3.99 mmol) in DMF (18 mL) and the stirred mixture heated at 115°C for 24 hours. Ether (100 mL) was added and washed with water (3 x 50 mL), brine (50 mL), dried (MgSO_4) and evaporated *in vacuo*. The residue was chromatographed on silica, eluting with a gradient of 50% ethyl acetate in isohexane, to give the desired compound (0.6 g)

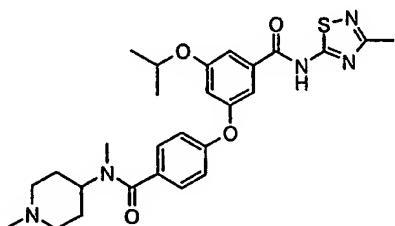
- 99 -

¹H NMR δ (CDCl₃): 1.35 (m, 9H), 3.78 (s, 3H), 4.36 (q, 2H), 4.58 (sept, 1H), 6.71 (m, 1H), 6.80 (m, 1H), 7.05 (m, 3H), 7.21 (m, 1H), 7.28 (m, 1H), 8.03 (d, 2H), 8.51 (s, 1H). m/z 424 (M+H)⁺

5 The preparation of 3-hydroxy-5-[(1-methylethyl)oxy]-N-(1-methyl-1*H*-pyrazol-3-yl)benzamide is described in Example 11.

Example 14: 3-[(1-Methylethyl)oxy]-5-[(4-{[methyl(1-methylpiperidin-4-

10 yl]amino]carbonyl}phenyl)oxy]-N-(3-methyl-1,2,4-thiadiazol-5-yl)benzamide

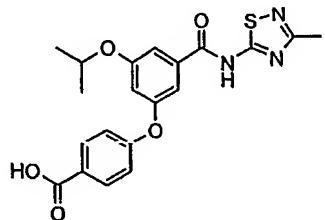


To a suspension of 4-[(3-[(1-methylethyl)oxy]-5-[(3-methyl-1,2,4-thiadiazol-5-yl)amino]carbonyl}phenyl)oxy]benzoic acid (250 mg, 0.61 mmol), HATU (480 mg, 1.27 mmol) and 1-methyl-4-(methylamino)piperidine (170 mg, 1.21 mmol) in DMF (4 mL), was
15 added diisopropylethylamine (0.44 mL, 2.24 mmol) and the mixture stirred at ambient temperature for 24 hours. Water (30 mL) was added and the mixture extracted with ethyl acetate (3 x 15 mL). The combined organic extracts were washed with brine, dried (MgSO₄), and evaporated to a residue which was chromatographed on silica, eluting with 10% methanol in ethyl acetate, to give the desired compound (52 mg).

20 ¹H NMR δ (CDCl₃): 1.29 (d, 6H), 1.59 (m, 2H), 1.80 (m, 4H), 2.12 (s, 3H), 2.37 (s, 3H), 2.80 (m, 6H), 4.70 (sept, 1H), 6.72 (m, 1H), 7.05 (d, 2H), 7.30 (m, 1H), 7.41 (d, 2H), 7.50 (s, 1H). m/z 524 (M+H)⁺

The required acid for Example 14 was prepared as described below:-

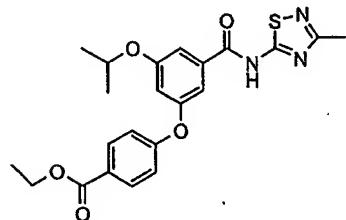
4-[(3-[(1-Methylethyl)oxy]-5-{{(3-methyl-1,2,4-thiadiazol-5-yl)amino]carbonyl}phenyl)oxy]benzoic acid



A solution of ethyl 4-[(3-[(1-methylethyl)oxy]-5-{{(3-methyl-1,2,4-thiadiazol-5-yl)amino]carbonyl}phenyl)oxy]benzoate (2.8 g, 6.35 mmol) in THF (105 mL) was added to a solution of lithium hydroxide monohydrate (1.33 g, 31.75 mmol) in water (53 mL). The mixture was stirred at ambient temperature for 48 hours and the THF removed *in vacuo*. The aqueous layer was acidified with 1M hydrochloric acid (30 mL), and the solid precipitate filtered off, washed with water and dried *in vacuo* to give the desired compound (2.6 g).

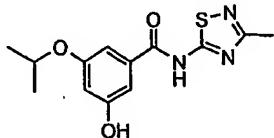
10 ^1H NMR δ (d_6 -DMSO): 1.31 (d, 6H), 2.45 (s, 3H), 4.72 (sept, 1H), 6.92 (m, 1H), 7.11 (d, 2H), 7.35 (m, 1H), 7.56 (m, 1H), 7.97 (d, 2H). m/z 414 ($M+\text{H}$)⁺

Ethyl 4-[(3-[(1-methylethyl)oxy]-5-{{(3-methyl-1,2,4-thiadiazol-5-yl)amino]carbonyl}phenyl)oxy]benzoate



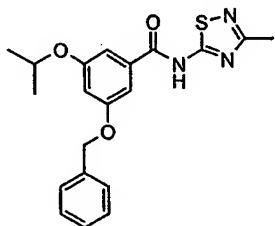
15 Potassium carbonate (9.38 g, 68 mmol) was added to a mixture of 3-hydroxy-5-[(1-methylethyl)oxy]-N-(3-methyl-1,2,4-thiadiazol-5-yl)benzamide (10 g, 34 mmol) and ethyl-4-fluorobenzoate (6.71 g, 41 mmol) in DMF (160 mL) and the stirred mixture heated at 115°C for 72 hours. Ether (300 mL) was added and washed with water (3 x 100 mL), brine (50 mL), dried (MgSO_4) and evaporated *in vacuo*. The residue was chromatographed on silica, eluting with a gradient of 5 to 50% ethyl acetate in isohexane, to give the desired compound (2.8 g)

20 ^1H NMR δ (d_6 -DMSO): 1.30 (m, 9H), 2.45 (s, 3H), 4.30 (q, 2H), 4.77 (sept, 1H), 6.95 (m, 1H), 7.15 (d, 2H), 7.36 (m, 1H), 7.55 (m, 1H), 7.99 (d, 2H), 13.38 (brs, 1H). m/z 442 ($M+\text{H}$)⁺

3-Hydroxy-5-[(1-methylethyl)oxy]-N-(3-methyl-1,2,4-thiadiazol-5-yl)benzamide

A solution of 3-[(1-methylethyl)oxy]-N-(3-methyl-1,2,4-thiadiazol-5-yl)-5-[(phenylmethyl)oxy]benzamide (33 g, 86 mmol), trifluoroacetic acid (160 mL) and thioanisole 5 (50.5 mL) was stirred at ambient temperature for 48 hours. The TFA was removed *in vacuo* and the residue poured into saturated sodium bicarbonate solution (300 mL) and extracted into ethyl acetate (twice). The combined organic extracts were washed with brine, dried (MgSO_4), filtered and the solvent removed *in vacuo*. The residue was triturated with DCM and washed with 5% ethyl acetate in isohexane to give the desired compound (12.8 g)

10 ^1H NMR δ (d_6 -DMSO): 1.31 (d, 6H), 2.51 (s, 3H), 4.67 (sept, 1H), 6.58 (s, 1H), 7.08 (s, 1H), 7.24 (s, 1H), 9.88 (s, 1H), 13.25 (brs, 1H). m/z 294 ($\text{M}+\text{H}$)⁺

3-[(1-Methylethyl)oxy]-N-(3-methyl-1,2,4-thiadiazol-5-yl)-5-[(phenylmethyl)oxy]benzamide

15 DMF (2 drops) was added to a solution of 3-[(1-methylethyl)oxy]-5-[(phenylmethyl)oxy]benzoic acid (29.6 g, 0.103 mol) and oxalyl chloride (10.78 mL, 0.12 mol) in DCM (500 mL). The mixture was stirred at ambient temperature for 4 hours and the DCM and excess oxalyl chloride removed *in vacuo*. The residual acid chloride was dissolved in DCM (220 mL) and added dropwise to 5-amino-3-methyl-1,2,4-thiadiazole (12.43 g, 0.108 mol) and triethylamine (30.34 mL, 0.216 mol) in DCM (220 mL), at 0°C. The reaction was allowed to warm and stirred at ambient temperature for 72 hours. The DCM was removed *in vacuo*, and the residue partitioned between ethyl acetate (400 mL) and 1N hydrochloric acid (200 mL). The ethyl acetate layer was washed sequentially with saturated aqueous sodium hydrogen carbonate (200 mL) and brine (100 mL), dried (MgSO_4) and concentrated *in vacuo*.

20 The residue was chromatographed on silica, eluting with a gradient of 20% ethyl acetate in isohexane, to give the desired compound (33 g)

25

¹H NMR δ (CDCl₃): 1.32 (d, 6H), 2.31 (s, 3H), 4.51 (sept, 1H), 5.05 (s, 2H), 6.74 (m, 1H), 7.03 (m, 1H), 7.10 (m, 1H), 7.38 (m, 5H), 11.48 (brs, 1H). m/z 384 (M+H)⁺

The preparation of 3-[(1-methylethyl)oxy]-5-[(phenylmethyl)oxy]benzoic acid is described in

5 Example 1.

BIOLOGICAL

Tests:

The biological effects of the compounds of formula (I) may be tested in the following way:

10

(1) Enzymatic activity

Enzymatic activity of recombinant human pancreatic GLK may be measured by incubating GLK, ATP and glucose. The rate of product (ie G-6-P) formation may be determined by coupling the assay to a G-6-P dehydrogenase, NADP/NADPH system and 15 measuring the linear increase with time of optical density at 340nm (Matschinsky et al 1993). Activation of GLK by compounds can be assessed using this assay in the presence or absence of GLKRP as described in Brocklehurst et al (Diabetes 2004, 53, 535-541).

Production of recombinant GLK and GLKRP

20 Human GLK and GLKRP cDNA was obtained by PCR from human pancreatic and hepatic mRNA respectively, using established techniques described in Sambrook J, Fritsch EF & Maniatis T, 1989. PCR primers were designed according to the GLK and GLKRP cDNA sequences shown in Tanizawa et al 1991 and Bonthon, D.T. et al 1994 (later corrected in Warner, J.P. 1995).

25 *Cloning in Bluescript II vectors*

GLK and GLKRP cDNA was cloned in E. coli using pBluescript II, (Short et al 1998) a recombinant cloning vector system similar to that employed by Yanisch-Perron C et al (1985), comprising a colEI-based replicon bearing a polylinker DNA fragment containing multiple unique restriction sites, flanked by bacteriophage T3 and T7 promoter sequences; a 30 filamentous phage origin of replication and an ampicillin drug resistance marker gene.

Transformations

E. Coli transformations were generally carried out by electroporation. 400 mL cultures of strains DH5a or BL21(DE3) were grown in L-broth to an OD 600 of 0.5 and harvested by centrifugation at 2,000g. The cells were washed twice in ice-cold deionised water,

5 resuspended in 1mL 10% glycerol and stored in aliquots at -70⁰C. Ligation mixes were desalted using Millipore V series™ membranes (0.0025mm) pore size). 40mL of cells were incubated with 1mL of ligation mix or plasmid DNA on ice for 10 minutes in 0.2cm electroporation cuvettes, and then pulsed using a Gene Pulser™ apparatus (BioRad) at 0.5kVcm⁻¹, 250mF. Transformants were selected on L-agar supplemented with tetracycline at

10 10mg/mL or ampicillin at 100mg/mL.

Expression

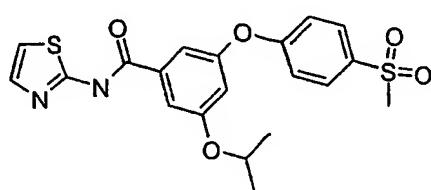
GLK was expressed from the vector pTB375NBSE in E.coli BL21 cells, producing a recombinant protein containing a 6-His tag immediately adjacent to the N-terminal methionine. Alternatively, another suitable vector is pET21(+)DNA, Novagen, Cat number 697703. The 6-His tag was used to allow purification of the recombinant protein on a column packed with nickel-nitrilotriacetic acid agarose purchased from Qiagen (cat no 30250).

GLKRP was expressed from the vector pFLAG CTC (IBI Kodak) in E.coli BL21 cells, producing a recombinant protein containing a C-terminal FLAG tag. The protein was purified 20 initially by DEAE Sepharose ion exchange followed by utilisation of the FLAG tag for final purification on an M2 anti-FLAG immunoaffinity column purchased from Sigma-Aldrich (cat no. A1205).

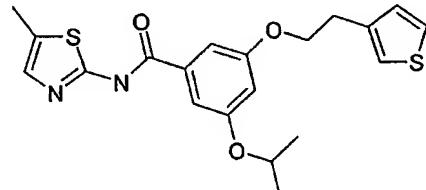
(2) Oral Glucose Tolerance Test (OGTT)

25 Oral glucose tolerance tests were done on conscious Zucker obese fa/fa rats (age 12-13 weeks or older) fed a high fat diet (45 % kcal fat) for at least two weeks prior to experimentation. The animals were fasted for 2 hours before use for experiments. A test compound or a vehicle was given orally 120 minutes before oral administration of a glucose solution at a dose of 2 g/kg body weight. Blood glucose levels were measured using a Accucheck glucometer from 30 tail bled samples taken at different time points before and after administration of glucose (time course of 60 minutes). A time curve of the blood glucose levels was generated and the area-under-the-curve (AUC) for 120 minutes was calculated (the time of glucose

administration being time zero). Percent inhibition is determined using the AUC in the vehicle-control group as zero percent inhibition.



Example 7c



Example II107

5 Compounds of the invention generally have an activating activity for glucokinase with an EC₅₀ of less than about 500nM. For example, Example 7c has an EC₅₀ of 50nM.

Example 7c and Example II107 in WO 03/015774 have broadly similar EC₅₀ values. However Example 7c has superior oral exposure and exhibits 18% OGTT activity at 10 mg/kg but Example II107 in WO 03/015774 is not active at 10 mg/kg.

10

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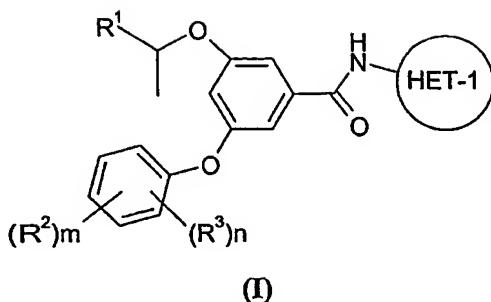
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December 2000

Claims:

1. A compound of Formula (I):



wherein:

R¹ is methyl;

R² is selected from -C(O)NR⁴R⁵, -SO₂NR⁴R⁵, -S(O)_pR⁴ and HET-2;

10 HET-1 is a 5- or 6-membered, C-linked heteroaryl ring containing a nitrogen atom in the 2-position and optionally 1 or 2 further ring heteroatoms independently selected from O, N and S; which ring is optionally substituted on an available carbon atom, or on a ring nitrogen atom provided it is not thereby quaternised, with 1 or 2 substituents independently selected from R⁶;

15 HET-2 is a 4-, 5- or 6-membered, C- or N-linked heterocyclyl ring containing 1, 2, 3 or 4 heteroatoms independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)-, and wherein a sulphur atom in the heterocyclic ring may optionally be oxidised to an S(O) or S(O)₂ group, which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁷;

20 R³ is selected from halo, fluoromethyl, difluoromethyl, trifluoromethyl, methyl, methoxy and cyano;

R⁴ is selected from hydrogen, (1-4C)alkyl [optionally substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted

25 with 1 group selected from R⁷) and HET-2;

R⁵ is hydrogen or (1-4C)alkyl;

or R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a heterocyclyl ring system as defined by HET-3;

R⁶ is independently selected from (1-4C)alkyl, halo, hydroxy(1-4C)alkyl, (1-4C)alkoxy(1-4C)alkyl, (1-4C)alkylS(O)p(1-4C)alkyl, amino(1-4C)alkyl, (1-4C)alkylamino(1-4C)alkyl, di(1-4C)alkylamino(1-4C)alkyl and HET-4;

R⁷ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkoxy(1-

5 5 10 15 20 25 30

HET-3 is an N-linked, 4, 5 or 6-membered, saturated or partially unsaturated heterocyclic ring, optionally containing 1 or 2 further heteroatoms (in addition to the linking N atom) independently selected from O, N and S, wherein a -CH₂- group can optionally be replaced by a -C(O)- and wherein a sulphur atom in the ring may optionally be oxidised to an S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an N-linked, 7 membered, saturated or partially unsaturated heterocyclic ring, optionally containing 1 further heteroatom (in addition to the linking N atom) independently selected from O, S and N, wherein a -CH₂- group can optionally be replaced by a -C(O)- group and wherein a sulphur atom in the ring may optionally be oxidised to an S(O) or S(O)₂ group; which ring is optionally substituted on an available carbon or nitrogen atom by 1 or 2 substituents independently selected from R⁸; or

HET-3 is an 6-10 membered bicyclic saturated or partially unsaturated heterocyclic ring, optionally containing 1 further nitrogen atom (in addition to the linking N atom), wherein a -CH₂- group can optionally be replaced by a -C(O)-; which ring is optionally substituted on an available carbon or nitrogen atom by 1 substituent selected from hydroxy and R³;

R⁸ is selected from -OR⁵, (1-4C)alkyl, -C(O)(1-4C)alkyl, -C(O)NR⁴R⁵, (1-4C)alkylamino, di(1-4C)alkylamino, HET-3 (wherein said ring is unsubstituted), (1-4C)alkoxy(1-4C)alkyl, hydroxy(1-4C)alkyl and -S(O)pR⁵;

HET-4 is a 5- or 6-membered, C- or N-linked unsubstituted heteroaryl ring containing 1, 2 or 3 ring heteroatoms independently selected from O, N and S;

p is (independently at each occurrence) 0, 1 or 2;

m is 0 or 1;

n is 0, 1 or 2;

provided that when m is 0, then n is 1 or 2;

or a salt, pro-drug or solvate thereof.

2. A compound of the formula (I) as claimed in Claim 1 or a salt, pro-drug or solvate thereof with the proviso that compounds exemplified in WO2004/076420, which would otherwise fall within the scope of this invention, are excluded.

5 3. A compound of the formula (I) as claimed in Claim 1 or Claim 2, or a salt, pro-drug or solvate thereof, wherein R¹ has the (S) configuration.

4. A compound of the formula (I) as claimed in Claim 1, Claim 2 or Claim 3, or a salt, pro-drug or solvate thereof, wherein HET-1 is a 5-membered ring.

10

5. A compound of the formula (I) as claimed in any one of Claims 1 to 4, or a salt, pro-drug or solvate thereof, wherein R² is selected from -C(O)NR⁴R⁵ and -SO₂NR⁴R⁵ and R⁴ and R⁵ together with the nitrogen atom to which they are attached may form a heterocyclic ring system as defined by HET-3.

15

6. A compound of the formula (I) as claimed in any one of Claims 1 to 5, or a salt, pro-drug or solvate thereof, wherein HET-3 is a 4- to 6-membered ring.

7. A compound of the formula (I) as claimed in Claim 1, Claim 2 or Claim 3, or a salt,
20 pro-drug or solvate thereof, wherein R² is selected from -C(O)NR⁴R⁵ and -SO₂NR⁴R⁵ and R⁴ is selected from (1-4C)alkyl [substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵], (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2.

25

8. A compound of the formula (I) as claimed in Claim 1, Claim 2 or Claim 3, or a salt, pro-drug or solvate thereof, wherein R² is -SO₂R⁴ and R⁴ is selected from (1-4C)alkyl [substituted by 1 or 2 substituents independently selected from HET-2, -OR⁵, -SO₂R⁵, (3-6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and -C(O)NR⁵R⁵], (3-30 6C)cycloalkyl (optionally substituted with 1 group selected from R⁷) and HET-2.

9. A compound of the formula (I) as claimed in Claim 1, Claim 2 or Claim 3, or a salt, pro-drug or solvate thereof, wherein R² is HET-2.

10. A pharmaceutical composition comprising a compound according to any one of
5 Claims 1 to 9, or a salt, pro-drug or solvate thereof, together with a pharmaceutically acceptable diluent or carrier.

11. A compound according to any one of Claims 1 to 9 or a pharmaceutically-acceptable salt, solvate or pro-drug thereof for use as a medicament.

10

12. A compound according to any one of Claims 1 to 9 for use in the preparation of a medicament for treatment of a disease mediated through GLK.

15

13. A compound according to any one of Claims 1 to 9 for use in the preparation of a medicament for treatment of type 2 diabetes.

14. A method of treating GLK mediated diseases by administering an effective amount of a compound of Formula (I) as claimed in any one of Claims 1 to 9 or salt, solvate or pro-drug thereof, to a mammal in need of such treatment.

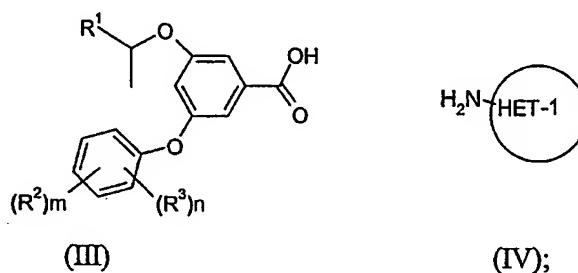
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15. The method of Claim 14 wherein the GLK mediated disease is type 2 diabetes.

25

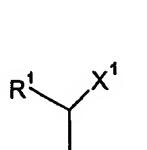
16. A process for the preparation of a compound of Formula (I) as claimed in any one of Claims 1 to 9, which comprises (wherein variables are as defined in Claim 1 unless otherwise stated):

(a) reaction of an acid of Formula (III) or activated derivative thereof with a compound of Formula (IV),

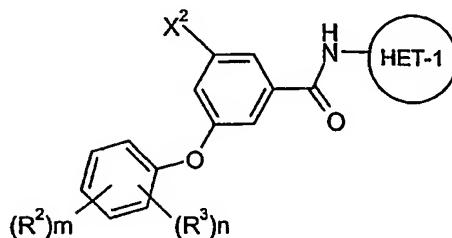


or

(b) reaction of a compound of Formula (V) with a compound of Formula (VI),



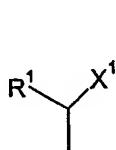
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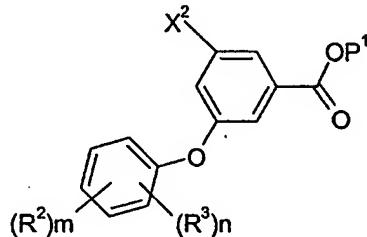
(VI)

5 wherein X¹ is a leaving group and X² is a hydroxyl group or X¹ is a hydroxyl group and X² is a leaving group;

[or by reaction with the intermediate ester Formula (VII), wherein P¹ is a protecting group followed by ester hydrolysis and amide formation];



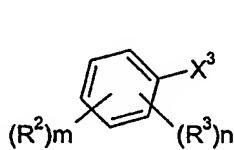
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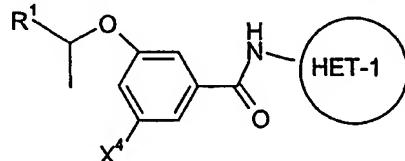
(VII)

or

(c) reaction of a compound of Formula (VIII) with a compound of Formula (IX)



(VIII)

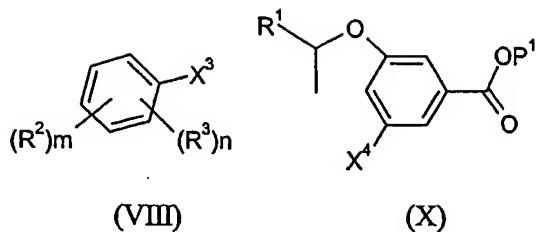


(IX)

15 wherein X^3 is a leaving group or an organometallic reagent and X^4 is a hydroxyl group or
 X^3 is a hydroxyl group and X^4 is a leaving group or an organometallic reagent;

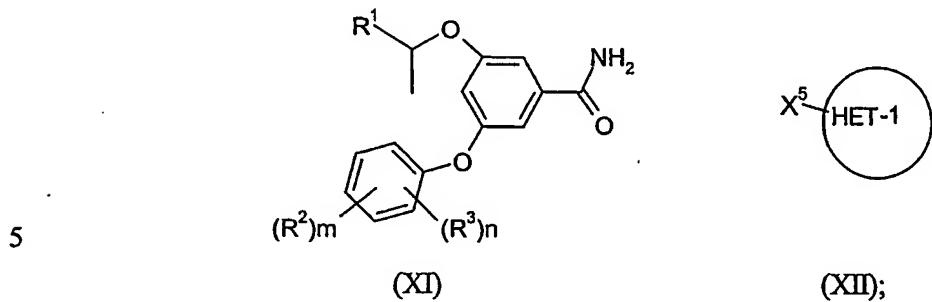
[or by reaction or (VIII) with the intermediate ester Formula (X), followed by ester hydrolysis and amide formation];

- 112 -



or

(d) reaction of a compound of Formula (XI) with a compound of Formula (XII),



wherein X⁵ is a leaving group;

and thereafter, if necessary:

i) converting a compound of Formula (I) into another compound of Formula (I);
10 ii) removing any protecting groups; and/or
iii) forming a salt, pro-drug or solvate.

17. A compound as exemplified herein, or a salt, pro-drug or solvate thereof.

INTERNATIONAL SEARCH REPORT

PCT/GB2005/000562

A. CLASSIFICATION OF SUBJECT MATTER					
IPC 7	C07D277/46	C07D417/12	C07D413/12	C07D405/12	C07D231/40
	C07D285/12	C07D213/75	C07D241/20	C07D263/48	C07D417/04
	C07D403/12	A61K31/426	A61P3/04	A61P3/06	

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, CHEM ABS Data, BEILSTEIN Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 03/015774 A (ASTRAZENECA AB; ASTRAZENECA UK LIMITED; BOYD, SCOTT; CAULKETT, PETER,) 27 February 2003 (2003-02-27) cited in the application Formulae (I), (IIb) claims 1-16	1-17
P,X	WO 2004/076420 A (BANYU PHARMACEUTICAL CO., LTD; IINO, TOMOHARU; HASHIMOTO, NORIAKI; NAK) 10 September 2004 (2004-09-10) cited in the application abstract; examples on pp. 111, 119, 120, 121, 123-125, 140, 153, 163, 164;	1-17

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the International filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the International filing date but later than the priority date claimed

- "T" later document published after the International filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "&" document member of the same patent family

Date of the actual completion of the international search

23 May 2005

Date of mailing of the International search report

31/05/2005

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INTERNATIONAL SEARCH REPORT

PCT/GB2005/000562

Box II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
Although claims 14,15 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. Claims Nos.: because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

PCT/GB2005/000562

Patent document cited in search report		Publication date		Patent family member(s)		Publication date
WO 03015774	A	27-02-2003	BR CA CN EP EP WO HU MX US	0212008 A 2457410 A1 1568185 A 1420784 A1 1529530 A1 03015774 A1 0401213 A2 PA04001500 A 2005080106 A1	28-09-2004 27-02-2003 19-01-2005 26-05-2004 11-05-2005 27-02-2003 28-12-2004 14-05-2004 14-04-2005	
WO 2004076420	A	10-09-2004	WO	2004076420 A1		10-09-2004